

<p style="text-align: center;">DECLARATION OF</p> <p style="text-align: center;">CAMERON JENNINGS, PH.D.</p> <p style="text-align: center;">UNDER 37 C.F.R. §1.132</p> <p>Address to: Commissioner for Patents P.O. Box 1450 Alexandria, VA 22313-1450</p>	Attorney Docket No.	29729/38914
	Confirmation No.	2874
	First Named Inventor	DUNN, ROSANNE D
	Application Number	10/590,690
	Filing Date	May 21, 2007
	Group Art Unit	1644
	Examiner Name	Schwadron, R.
	Title:	"TARGET FOR B-CELL DISORDERS "

Sir:

1. I, Cameron Jennings, PhD, declare and say I am a resident of 1 Janice Street, Warners Bay, New South Wales 2282, Australia.
2. I am an employee of Energy and Management Services Pty Ltd. My previous scientific employment consists of Immune System Therapeutics Ltd (Sydney), Harvard University (Boston) and La Trobe University (Melbourne). Details of my career as well as publications may be found in my curriculum vitae (Exhibit A).
3. I understand that US Patent Application No. 10/590,690 (referred to hereafter as the "Patent Application") is assigned to Immune System Therapeutics Limited. I presently own options which would allow me to purchase shares in Immune System Therapeutics Limited should the company become publicly listed.
4. I have been asked by FB Rice, Patent Attorneys for Immune System Therapeutics Limited, to provide an independent opinion on the state of knowledge surrounding kappa and lambda light chains and on the invention described in the Patent Application. I have been asked in particular to comment on the obviousness rejections set out in the Office Action dated 11 September 2009 issued in connection with the Patent Application.
5. I have read the Patent Application and have reviewed the claims that I understand are presently being considered by the United States Patent and Trademark Office. I understand that the claimed invention (referred to hereafter as the "Invention") relates to methods of treating B-cell disorders by administering an antibody or ligand that specifically binds to lambda myeloma antigen (LMA). The importance of this invention is that it provides, for the first time, LMA as a target that is selective for tumor B-cells

expressing LMA. Antibodies and ligands that bind LMA do not target normal B-cells which express lambda light chain in the context of intact immunoglobulin.

6. I have considerable experience in the technical field of the claimed invention. I have over 3 years of experience in research involving immunoglobulin light chains and membrane-bound proteins. Prior to this time, my research consisted of a senior post-doctoral fellowship at Harvard University researching membrane bound malaria vaccine candidates and a PhD that was co-supervised by Professor Marilyn Anderson (Biochemistry Department at La Trobe University) and Professor David Craik (Institute of Molecular Bioscience at The University of Queensland).
7. I have reviewed the Office Action dated 11 September 2009 in relation to the above-referenced Patent Application. I understand that the Patent Office has taken the position that the Invention would be obvious to a person skilled in the art in light of Uhr *et al.* (US 7,792,447), Raison *et al.* (WO 03/004056) and Abe *et al.* (1993).
8. Uhr *et al.* are cited as disclosing antibodies that bind intact immunoglobulin associated lambda light chain on tumor cells. Raison *et al.* are cited as disclosing that malignant cells express both kappa and lambda light chains, that free kappa light chain is expressed on the cell surface of kappa light chain expressing myeloma cells and that antibodies which bind free kappa light chain can be used to treat tumors. Abe *et al.* are cited as disclosing antibodies that bind free light chain and not intact immunoglobulin.
9. It is my understanding that the Patent Application and the cited publications are to be viewed from the perspective of one of ordinary skill in the art in the relevant field (a "Skilled Person") at the time of filing of the Patent Application in question. I have been asked to consider this time to be the period around or before 27 February 2004 ("the Relevant Period"). I would expect a Skilled Person in the field of antibody therapy during the Relevant Period to have been represented by a scientist with a PhD degree in Biochemistry and/or at least 3 to 5 years experience in the field of Biochemistry, or an educational background at the same degree level in a related field and equivalent level of experience.
10. I am very familiar with the technical field of the claimed invention. I am qualified to analyze literature in this field and to provide my opinion as to what literature in this field discloses or suggests to the Skilled Person at the Relevant Period.

11. By the Relevant Period I had attained at least the level of such a Skilled Person, and further in view of my qualifications discussed above, I believe that I am qualified by training and experience to address what a Skilled Person would have understood from reading the Patent Application and the cited publications.
12. The Examiner's objection appears to be based on the premise that kappa and lambda light chains are the two known alleles of immunoglobulin light chain and therefore once free kappa light chain had been found expressed on the surface of myeloma cells, it would have been expected that free lambda light chains would also be expressed on the surface of malignant B cells.
13. In my view, it is not correct to assume that the expression and localisation of the kappa light chain in myeloma cells is in any way predictive of the expression and localisation of lambda light chains in malignant B cells.
14. Immunoglobulins are composed of two chains, light and heavy, which form a functional heterodimer. The light chain molecules are of two protein families, "kappa" and "lambda". The kappa and Lambda gene loci are located on chromosomes 2 and 22 respectively and differ in their number of variable (V), joining (J) and constant (C) genes as well as their general arrangement. For example, there is a single C kappa gene (Zimmer *et al.*, 1990) while there are at least 4 functional C lambda genes (Dariavach *et al.*, 1987). While both kappa and lambda light chains maintain a conserved role when present in intact immunoglobulin, no normal biological function has been attributed to light chains alone.
15. Although kappa and lambda light chains both comprise a conserved (C) and variable (V) domain (within the respective families) and they both complete a multimeric immunoglobulin, the proteins share minimal sequence identity within their C domains (Kabat *et al.*, 1975). Furthermore, the variable domains of the kappa and lambda light chains are derived from genetic recombination events and the recombined genes also undergo the process of somatic hypermutation to give rise to antigen binding moieties (in the context of the whole immunoglobulin). As such the V regions vary in their sequence as well.
16. Kappa and lambda V domains have a typical Immunoglobulin fold, which is also referred to as the immunoglobulin beta sandwich, consisting of two antiparallel β -barrel sheets.

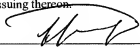
Although both kappa and lambda V domains comprise this typical Immunoglobulin fold, analysis of the crystal structure of light chains demonstrated that the C domains of lambda light chain and gamma heavy chain were structurally amenable to the formation of cross-molecule beta structures, but kappa light chains were not (Edmundson and Borrebaeck, 1998). Thus, it was known that there were differences in the structural interactions between lambda light chains and immunoglobulin heavy when compared to the interactions of kappa light chain and heavy chain.

17. The immunoglobulin beta sandwich, referred to above, contains seven beta strands which form the sandwich of two beta sheets. This structural motif is found on a vast array of protein sub-families with diverse biological activities and sub-cellular locations (for a complete set of proteins in the SCOP data base, see (Murzin *et al.*, 1995; available from <http://scop.mrc-lmb.cam.ac.uk/scop/data/scop.b.c.b.html>). Proteins that fall within this structural family include Myelin oligodendrocyte glycoprotein (MOG), the T-cell antigen receptor and TREM-1 (triggering receptor expressed on myeloid cells) to name a few.
18. As further evidence of differences in structure of kappa and lambda light chain, the *Peptostreptococcus magnus* protein L binds kappa light chains but does not bind lambda light chains (Graille *et al.*, 2001). This again suggests variation in sequence and local structure of the kappa and lambda light chains
19. The differences in the structures of the kappa and lambda light chains is further highlighted by observations indicating that lambda antibodies are found in two-thirds of light chain Amyloidosis cases, whereas kappa light chains mediate greater than 85% of Light-Chain Deposition Disease (LCDD). Thus, the light chain composition appears to confer different pathologies to kappa or lambda light chains (Solomon and Weiss, 1995).
20. The difference in the structure of amyloid fibrils (fibrillar) and the LCDD deposits (amorphous) also reflects the difference in the general structure of kappa and lambda light chains (Khurana *et al.*, 2001). Moreover, lambda light chains exist predominantly as dimers while kappa light chains are mostly present as monomers (Solomon and Weiss, 1995). As a consequence, the character and rate of the catabolic processes that are involved in the clearance of kappa and lambda light chains are different. It has been suggested that this phenomenon contributes to the predominance of lambda light chains in amyloidosis (Solomon and Weiss, 1995).

21. There exists three broad categories of surface expressed proteins: i) integrally associated proteins possessing hydrophobic surfaces that readily interact with the acyl core of the bilayer; ii) membrane proteins that are covalently attached to certain phospholipids; and iii) peripheral proteins that associate with the membrane through charge-charge electrostatic interactions (Lodish et al. (2000) Molecular and Cellular Biology 4th Ed., Section 3.4) WH Freeman, New York). Examination of the sequence of kappa and lambda light chains does not reveal candidate residues suitable for covalent attachment to the membrane. Thus, it can be surmised that the membrane interaction of either kappa or lambda light chains would occur via hydrophobic and/or electrostatic interactions.
22. Differences in the primary sequence of kappa and lambda light chains affects not only the charge of the exposed side chains of the proteins but also their predominant presence as either monomers or dimers respectively. Thus, in view of the differences in primary sequence of kappa and lambda light chains, which in turn affects the presence/absence of exposed hydrophobic surfaces, and the lack of a membrane targeting sequence in these proteins, the expression of either of these two proteins on the cell surface could not be predicted. Thus, it could not have been predicted by analysis of the structure or sequence of kappa or lambda light chains that either of these proteins would associate with the membrane of malignant B cells.
23. In my opinion, therefore, there is nothing in the cited prior art to suggest that a Skilled Person could have predicted that free lambda light chain would be associated with the membrane of tumor B-cells. There was no suggestion nor motivation, therefore, around February 2004 for a Skilled Person to investigate free lambda light chains as a potential therapeutic target on tumor B-cells.
24. I hereby declare that all statements made herein of my own knowledge are true and that all statements made on information and belief are believed to be true; and further that these statements were made with the knowledge that willful false statements and the like so made are punishable by fine or imprisonment, or both, under § 1001 of Title XVIII of the United States Code, and that such willful false statements may jeopardize the validity of the Patent Application or any patent issuing thereon.

2/3/2012

Date



Name: Cameron Jennings, PhD

EXHIBIT A

Dr Cameron Victor Jennings

1 Janice Street, Warners Bay NSW 2282

Australia

+61430095522

cam_jennings@hotmail.com

EDUCATION:

1. Post-Graduate Diploma in Energy Studies (Awarded in 2010)

Murdoch University, School of Engineering and Energy
Perth, Australia

2. Doctor of Philosophy (awarded in 2003)

La Trobe University, Department of Biochemistry
Melbourne Australia.

3. Bachelor of Science (First Class Honours-- awarded in 1999)

La Trobe University
Melbourne Australia.

WORK EXPERIENCE:

1. Senior Scientist/Project Leader (July 2007 – Present)

Immune System Therapeutics LTD. Sydney Australia.

Role: Supervise and drive projects that increase the intellectual property profile of Immune System Therapeutics Ltd.

2. Scientist (2006-July 2007)

Immune System Therapeutics LTD. Sydney Australia.

Role: Drive projects that increase the intellectual property profile of the company

3. Senior Post-Doctoral Research Fellow (2003-2006)

**Harvard University, School of Public Health,
Department of Immunology and Infectious Diseases, Boston USA.**

Role: Perform and supervise research into Malaria vaccine candidates

4. Laboratory class teaching assistant - Undergraduate Biochemistry (1999-2001)

La Trobe University, Department of Biochemistry. Melbourne, Australia.

Role: Supervise undergraduate students in practical classes

5. Research Assistant, Phillips Laboratory (1997-1999).

La Trobe University, Department of Biochemistry. Melbourne, Australia.

Role: Support research into potential anti-cancer compounds

OTHER ACADEMIC TRAINING:

1. Environmental Auditor Certification Workshop (July, 2009)

Thomson - Reuters

Aim: Training and certification for auditing environmental management systems

2. Environmental Management Systems Workshop (July, 2009)

Thomson - Reuters

Aim: Additional training in environmental management systems and auditing

3. Integrated Management Systems Workshop (July, 2009)

Thomson - Reuters

Aim: Training in integrating environmental management systems into existing business structures

4. NSW Enterprise Workshop (Winter Program 2008)

Aim: To develop and defend a business plan, including an executive summary, marketing, operations, project management and a financial plan

ADDITIONAL ACHIEVEMENTS:

- Won a **business plan award** at the NSW Enterprise Workshop (Best Intellectual Property Plan, shared award with 2 other team members) 2008
- Supervised an honours student who received the **Deans Merit Award** at the University of Technology, Sydney 2008
- Jennings et al., 2005, was among the 20 most cited articles (position 11) of 2005, January – June.
- Awarded a monetary prize for the **best oral seminar** at the inaugural Melbourne Protein Group Symposium. 2002.
- Received a **student bursary** to attend the Forth Australian Peptide Conference (Lindeman Island, Queensland, Australia). 2001.
- **President of the La Trobe University Biochemistry Society.** 2000
- Accepted into the **Golden Key Club** for Scholastic achievements. 1997.
- Accepted onto the **Deans List** for Scholastic achievements (La Trobe University, Melbourne, Australia. 1997.

PATENTS:

Novel nucleic acid molecules

International application WO 0134829 filed 25th November 2000.

Inventors: Craik, D.J., Anderson, M.A. and Jennings, C.V.

Several Provisional patent applications written and filed

INVITED SEMINARS/ORAL PRESENTATIONS:

- **Institute for the Biotechnology or Infectious Diseases (University of Technology Sydney) 2007**
Jennings C.V., Ahouidi A.D., Bei A., Sarr O., Ndir O., Wirth D., Mboup S., Duraisingh M.T
- **Biological Sciences and Public Health Retreat (Harvard University) 2005**
Jennings C.V., Ahouidi A.D., Bei A., Sarr O., Ndir O., Wirth D., Mboup S., Duraisingh M.T.
- **Biological Sciences and Public Health Retreat (Harvard University) 2003**
Jennings, C.V., Whitehurst, N., Desimone, A., Duraisingh, M.T.
- **PhD Qualification Seminar 2002**
Jennings C.V. Craik D.J., Anderson M.A
- **Inaugural Melbourne Protein Group Symposium 2002**
Jennings C.V., West J.A., Craik D.J., Anderson M.A.
- **PhD Conversion Seminar 2001**
Jennings C.V. Craik D.J., Anderson M.A

RESEARCH PUBLICATIONS:

Bushkin GG, Ratner DM, Cui J, Banerjee S, Duraisingh MT, **Jennings CV**, Dvorin JD, Gubbels MJ, Robertson SD, Steffen M, O'Keefe BR, Robbins PW, Samuelson J. Eukaryot Cell. 2009 Oct 16.

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Mol Microbiol. 2009 Epub. Apr 14

* Joint first authors

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Mulvenna, J.
International Journal of Peptide Research and Therapeutics, 2001 May;8(3-5): 119-128.

Transposition of human immunoglobulin V_κ genes within the same chromosome and the mechanism of their amplification

F.-J. Zimmer, H. Hameister¹, H. Schek and H. G. Zachau

Institut für Physiologische Chemie, Physikalische Biochemie und Zellbiologie der Universität München, 8000 München 2 and ¹Abteilung Klinische Genetik der Universität Ulm, 7900 Ulm, FRG

Communicated by H. G. Zachau

The variable, joining and constant gene segments of the human immunoglobulin κ locus (V_κ, J_κ and C_κ) are located on the short arm of chromosome 2 at 2p11–2p12. Here we describe a cluster of 11 V_κ genes on the long arm of chromosome 2 at 2cen–q11. By pulsed-field gel electrophoresis, cosmid cloning and DNA sequencing the cluster was shown to consist of four amplified units (amplicons). The amplicons, each 110–160 kb in size, are organized within 650 kb as an array of inverted repeats with short stretches of non-amplified DNA in between. Cloning and sequencing of three different joints between amplified and non-amplified DNA revealed the existence of parts of Alu repeats at each of the analysed joints. It is suggested that during evolution a group of five V_κ genes was transposed from the short to the long arm of chromosome 2 by a pericentric inversion. Three of the five V_κ genes were then amplified in two subsequent steps to yield the structure found in the majority of the present day population. The possible relation of this structure to a pericentric inversion of chromosome 2 that is seen cytogenetically in a small fraction of today's population is discussed.

Key words: Alu repeats/amplification/immunoglobulin V_κ genes/transposition

Introduction

Amplification and transposition of genes play an important role in the formation of multigene families during evolution (for review, see Maeda and Smithies, 1986). In the case of the human gene family coding for the variable regions of the immunoglobulin light chains of the κ type (V_κ), putative transpositions of V_κ genes led to the formation of a 'mixed' gene cluster, in which genes of different subgroups are interdigitated (Pech and Zachau, 1984). A subsequent duplication led to the generation of a second copy of a large part of the V_κ gene cluster (for reviews, see Zachau, 1989, 1990). The κ locus contains ~70 V_κ genes (for reviews, see Zachau, 1989, 1990) and is located on the short arm of chromosome 2, at 2p12 (Malcolm *et al.*, 1982). In addition, V_κ genes have been found on the chromosomes 1, 22 (Lötscher *et al.*, 1986, 1988) and other chromosomes (Straubinger *et al.*, 1988). These genes are called orphans in analogy to histone and ribosomal RNA genes found outside of their respective gene clusters (Childs *et al.*, 1981).

We have previously reported the structure of two con-

tiguous cloned regions (contigs), called Wa and Wb, with a total of nine V_κ genes (Pohlentz *et al.*, 1987). The two W contigs have been assigned to chromosome 2 (Lötscher *et al.*, 1988) and were thought to be part of the κ locus. However, we did not succeed in linking them to the cloned parts of the κ locus by chromosomal walking or by pulsed-field gel electrophoresis (PFG). In the course of the PFG experiments, a third W contig was detected. The three contigs are present in the genomes of all individuals so far analysed. The observation that the W contigs are located on chromosome 2, yet are not part of the κ locus, prompted us to analyse their genomic organization and chromosomal location in more detail.

Results

Characterization of the W contigs

We previously described the characterization of the two contigs Wa and Wb (Pohlentz *et al.*, 1987). Here, we report the isolation and characterization of two sets of cosmid clones, one extending the contig Wb and one representing a new W contig, termed Wc.

In the course of chromosomal walking experiments, the cosmid libraries III (Pohlentz *et al.*, 1987) and V (Lorenz, 1989) were screened with the W-specific clone m654-1 (Pohlentz *et al.*, 1987); 25 cosmid clones were isolated. Five of them were derived from Wa without extending the known contig. The other 20 clones belonged to Wb with some of them extending the contig. Two additional V_κ genes were found in the extending cosmids. A map of the W contigs with some representative cosmid clones is shown in Figure 1. A description of all clones can be found in Zimmer (1989).

The existence of a third contig, Wc, was demonstrated on PFG blots that had been hybridized with the Wa-derived probe m167-1 (Pohlentz *et al.*, 1987; see also Figure 1). In subsequent restriction mapping of cosmid clones that had been isolated previously with m167-1, but not fully characterized (Pohlentz, 1986), three clones matching neither Wa nor Wb were identified. Wc was shown to be a third independent contig and not an allelic variant of Wa or Wb by demonstrating the existence of Wa-, Wb- and Wc-characteristic fragments in the genomes of all 20 unrelated individuals tested (Zimmer, 1989; data not shown here). The map of Wc is included in Figure 1.

We use the term amplified unit or amplicon for those parts of the contigs Wa, Wb and Wc, that are homologous to each other. As Wb contains two amplified units, a minimum of four W amplicons (named I–IV; Figure 1) exist within the human genome.

How many V_κ gene containing W amplicons exist in the human genome?

To test whether more than three V_κ gene-containing amplicons exist, we estimated their copy number relative to C_κ, which is known to be single copy, by quantitation

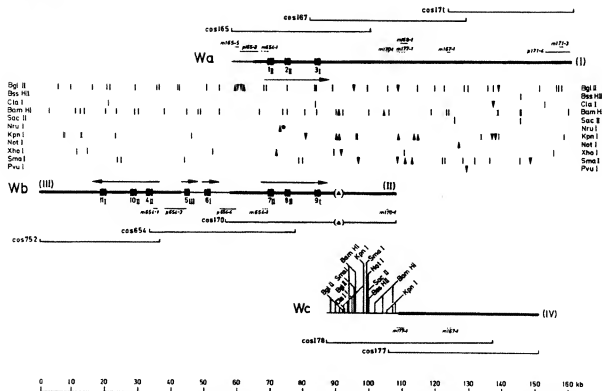


Fig. 1. Restriction maps of the genomic contigs Wa, Wb and Wc. The maps were derived from cosmid clones described in Pohlentz *et al.* (1987) and Zimmer (1989). For each contig only some representative cosmid clones are shown. The maps are aligned to demonstrate the homology between the contigs. Positions of restriction sites that are identical in two contigs or occur in contigs without counterpart are marked by vertical bars.

Differences in restriction sites between two contigs are symbolized by arrows pointing to the respective map positions. In the map of Wc, restriction sites are given separately for the part that shows no homology to Wb or Wc. The scale (kb) applies to all contigs. A deletion of 1 kb in Wb at map position 92 kb is marked by a triangle. V_{α} genes are drawn as filled boxes; the subgroup designations are indicated (I–III). Plasmid and M13 subclones are shown as horizontal bars above the restriction map of Wa and underneath those of Wb and Wc. The subclones m654-1, m655-5 and m167-1 are described in Pohlentz *et al.* (1987), the remaining ones in Zimmer (1989). Amplified units are marked by fat lines and are numbered (I–IV). Clones derived from one amplicon hybridize to the corresponding position within other amplicons; these positions are marked by dotted lines. Arrows at the maps of Wa and Wb indicate the transcriptional orientation of the V_{α} genes. A dot marks a *Nru*I site at map position 72 kb, which is present only in some alleles of Wa.

of Southern blot hybridizations. We chose a clone which hybridizes to a position close to one of the amplified V_{α} genes (m654-1) and linked it to a fragment derived from the C_{α} region (I-1). The construct m654-1/I-1 is shown in Figure 2a and the blot hybridizations are in Figure 2b. The copy number of the W contigs is estimated as the multiple of the C_{α} signal (Table I). From the calculated W/ C_{α} ratios it is very likely that no further amplicons hybridizing with m654-1 exist. The procedure of copy number determination is similar to the one of Meindl (1990) who estimated the overall number of V_{α} genes including W-type V_{α} genes in the DNA of individual AF, and arrived at data compatible with three copies of the W amplicon and one copy of the non-amplified gene W6. The cosmid clones constituting the W contigs of Figure 1 are derived from the DNA of individual N and individual St. Since the blot hybridizations of restriction nuclease digests of the DNAs AF, N, St and PC-3 (see below) show no significant differences in the regions of the W-type V_{α} genes (Meindl, 1990) it is likely that the number of V_{α} gene-containing W amplicons is three. The amplified unit within Wc is not detected by hybridization with m654-1/I-1 since it does not reach into the V_{α} gene-containing regions of the W amplicons (Figure 1). To identify other possibly existing truncated

amplicons and to elucidate the genomic organization of the known amplicons, we established a long-range map of all regions hybridizing with W-specific probes by PFG.

Long-range map of the W regions

The analysis of the organization of the W region was carried out with DNA from the prostate carcinoma cell line PC-3 (Kaighn *et al.*, 1979), as PC-3 DNA seems to contain more unmethylated restriction sites than DNA from other sources (Lorenz *et al.*, 1987), allowing the use of methylation sensitive restriction nucleases for long-range mapping.

A remarkable feature of the PFG blots is that all W region probes tested hybridize only to two *Not*I fragments, 250 and 600 kb in size. V_{α} gene probes hybridize mainly with two additional *Not*I fragments which contain the α locus (Lorenz *et al.*, 1987). The identification of *Not*I sites in cosmid clones of Wa and Wc, which are also present in PC-3 DNA (*Not*I sites at map positions 550 and 1150 kb of Figure 3), has been useful in the construction of the long-range map.

The PFG studies did not reveal any evidence for the existence of additional W regions. According to the PFG experiments, the four W amplicons are organized as an array of large inverted repeats.

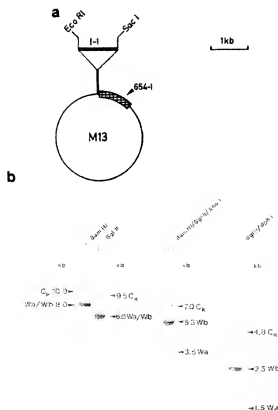


Fig. 2. Determination of the number of W contigs. In (a) the construct m654-1/I-1, which was used for the hybridizations, is shown. A 1.1 kb *EcoRI*–*SacI* fragment from pl-1, which is unique for the C_h region (Klobeck *et al.*, 1984), has been cloned into the W-specific probe m654-1. Southern blot analyses with m654-1/I-1 are shown in (b). Each lane contains 10 µg of placenta DNA digested with the indicated restriction nucleases. The W and C_h derived fragments are assigned to the respective bands, the sizes of which are given in kb. For the quantitative evaluation see Table I.

Cloning and sequence analyses of joints between amplified and non-amplified DNA

Four joints between amplified and non-amplified DNA can be identified within the W regions by comparing the respective restriction maps (transitions between bars and lines in Figure 1). By cloning joint-containing fragments into plasmid (p165-3, p654-3, p654-4) or M13 vectors (m177-1, m168-1) and comparing the fine maps, fragments suitable for the sequence analyses of the joints were identified and subcloned.

The break-off in homology between Wa and Wb was localized by aligning the map of Wa with that of the 5' amplified unit of Wb (amplicon III) and comparing the restriction maps of p165-3 and p654-3. The joints of the amplicons II and III in Wb were localized in a similar manner by comparing the maps of p654-3 and p654-4. To localize the Wc joint, the maps of m168-1 and m177-1 were aligned (Figure 1). The sequences of the analysed joints are shown in Figure 4. It is evident from these data that Alu repeats

Table I. Determination of the number of V_h gene-containing W amplicons

Restriction digests	Size of bands ^a (kb)	Region ^a	Counts/min ^b	Copy number ^c
<i>Bam</i> HI	10.0	C _h	14 ± 10	1.0
	8.0	Wa + Wb	47 ± 9	3.5 (3–4)
<i>Bgl</i> II	9.5	C _h	21 ± 10	1.0
	6.6	Wa + Wb	58 ± 9	2.8 (3)
<i>Bam</i> HI/ <i>Bgl</i> II	7.0	C _h	43 ± 7	1.0
<i>Xho</i> I	5.3	Wb	87 ± 7	2.0 (2)
	3.5	Wa	32 ± 8	0.7 (1)
<i>Bgl</i> II/ <i>Sph</i> I	4.8	C _h	40 ± 7	1.0
	2.5	Wb	80 ± 7	2.0 (2)
	1.5	Wa	35 ± 8	0.9 (1)

^aThe assignment of bands in the digests to certain contigs is seen in Figure 2b.

^bThe values are corrected for blank values as described in Materials and Methods. The standard deviations are indicated.

^cThe copy number for C_h is taken as 1.0. The W/C_h ratios are calculated by dividing the c.p.m. of the W containing bands by the one for C_h; the resulting numbers of W copies are given in parentheses.

played a role in the recombination processes, which led to the formation of the novel joints.

The W contigs reside on the long arm of chromosome 2

As we have not been able to link by chromosomal walking or PFG the W contigs to the *x* locus, we tried to determine the chromosomal location of W by analysing somatic cell hybrids.

In one set of experiments, mouse–human hybrid cell lines, which contain only parts of human chromosome 2, were analysed with probes specific for the *x* locus or the W regions. In one of the lines, J14-2L (Erikson *et al.*, 1983), only that part of the short arm of chromosome 2 is present that comprises C_h to the telomere (2p12–tel). DNA of this cell line hybridizes only with a C_h-specific probe but with neither probes derived from the V_h gene-containing parts of the locus nor the W region-specific probe m654-1 (Zimmer, 1989).

A second analysed cell line, RRP5-3 (Shiloh *et al.*, 1985), represents the almost complementary situation to J14-2L, as RRP5-3 contains a 2p⁺ chromosome, i.e. the long arm and only that part of the short arm between the centromere and the *x* locus in 2p12. DNA of this cell line does not hybridize with any of the single-copy probes specific for the *x* locus; however, it gives a strong signal with m654-1 (Zimmer, 1989). According to these data, the W contigs are located either on the short arm of chromosome 2, between the *x* locus and the centromere, or on the long arm.

A more precise localization of the W regions was achieved by *in situ* hybridizations; these results are shown in Figure 5. The *x* locus specific probe pC-2 (Klobeck *et al.*, 1984) maps to the short arm of chromosome 2 in the region 2p11–2p12 in accordance with the location of the *x* locus (Malcolm *et al.*, 1982; McBride *et al.*, 1982). The W-derived cosmid clones cos178, cos165 (Figure 1) and cos140 (Pohlentz *et al.*, 1987) as well as the subclone m654-1 (data not shown) map to a region close to the centromere on the long arm of

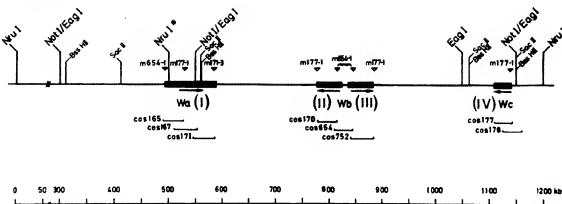


Fig. 3. Long range map of the W contigs. The map was constructed on the basis of numerous PFG blots, most of which were hybridized consecutively with different probes in order to find out whether certain fragments are recognized by more than one probe [a list of fragments hybridizing with W-specific probes can be found in Zimmer (1989)]. The shown orientation of Wb is arbitrary, as there is no cloned restriction site within Wb which would allow one to determine its orientation. Wa and Wc were oriented by identification of restriction sites for *NotI* and *NruI* in cosmids of the Wa contig, and of a *NotI* site in cosmids of the Wc contig. These sites are also present in DNA of PC-3 which is used in the PFG experiments (map positions 500, 550 and 1150 kb). The distance between the Wa and Wc contigs is defined by the 600 kb *NotI* fragment which hybridized with m171-3 and m177-1. The contig Wb is placed in the centre of this fragment since we assume about equal sizes of the amplicons (see Discussion). The 500 and 700 kb *NruI* fragments are defined by hybridization with m654-1, m171-3 and m177-1. The amplicons I–IV are shown as black bars. Arrows indicate the amplicon orientation based on the transcriptional orientation of *V_h* genes (I–III); the orientation of amplicon IV is based on its restriction map which is homologous to that of amplicon I (Figure 1). Filled and open triangles mark the regions to which the indicated probes hybridize. Open symbols indicate that the hybridizing region is not represented by cosmid clones. The terminal part of the map without known restriction sites is shortened (—//—). The analysed cell line PC-3 (Kaighn *et al.*, 1979) is heterozygous for the *NruI* site marked by a rhombus. The linking of Wa to Wb and Wb to Wc is based on those restriction fragments marked with larger letters [for details see Zimmer (1989)].

chromosome 2 (2cen–q11). This position is obviously distinct from the *x* locus.

Discussion

Size, copy number and organization of the W region amplicons

According to the PFG experiments and the copy number determinations, four amplified units hybridizing with W region-specific probes reside within a DNA stretch of 650 kb and are organized as an array of inverted repeats. The cloned parts of the amplicons I–IV are 50–100 kb each (Figure 1). If the length of non-amplified DNA between the amplicons I and II and between III and IV (Figure 3) is similar to that between the amplicons II and III, the average size of the amplicons is in the range 110–160 kb. The chromosomal organization of the W amplicons, i.e. variable amplicon size, head-to-head or tail-to-tail orientation of the amplified units and stretches of non-amplified DNA in between two amplicons, is very similar to the organization of, for example, the amplified genes of dihydrofolate reductase (DHFR), CAD, *c-myc* and adenylate deaminase (AMPD) (DHFR, Ma *et al.*, 1988; Hearle and Latt, 1989; CAD, Ardeshrir *et al.*, 1983; Ford and Fried, 1986; *c-myc*, Ford and Fried, 1986; AMPD, Hyrien *et al.*, 1988).

One major difference between the organization of the W regions and the amplified units reported in the literature is the low copy number and a rather small amplicon size. The amplicons described so far exist either in a low copy number, which is the case after first step selection, but have a size up to 10 000 kb, found for CAD amplicons (Giulotto *et al.*, 1986) or the size is in the range of that of the W region amplicons, but the copy number reaches a value up to 2600

copies per cell, as has been described for AMPD amplicons (Yeung *et al.*, 1983). Whether these differences reflect a different mechanism of amplification responsible for the generation of the W regions is discussed in a following section.

Novel joints are formed within Alu repeats

Relatively few studies define the DNA sequences at the sites of recombination associated with amplification (for review, see Stark *et al.*, 1989). The sequence analyses of novel joints revealed the existence of partial Alu repeats. This finding makes it very likely that the repeats played a role either during the amplification process itself or in recombinations, which are necessary to resolve aberrant replication bubbles into a linear array of amplified units. Depending on the assumed mechanism of amplification (for review, see Stark *et al.*, 1989), one can imagine several ways in which Alu repeats participate in the amplification process. In the context of strand switch models (Nalbantoglu and Meuth, 1986; Hyrien *et al.*, 1988), Alu repeats could serve as those sequences which promote the strand switch by the DNA polymerase within the replication fork. According to such a model, one would expect to find partial Alu repeats at the amplification joints if the strand switch event did not occur at identical positions on the leading and lagging strand.

In the course of recombination, which is always associated with DNA amplification, Alu repeats might serve as target sites for the recombinations. The involvement of Alu repeats in genomic recombinations has been reported frequently (see literature cited in Hyrien *et al.*, 1987).

Although we cannot decide at which step during the generation of the W amplicons the Alu repeats played a role, these repetitive sequence elements seemed to be important for the amplification of a W region precursor.

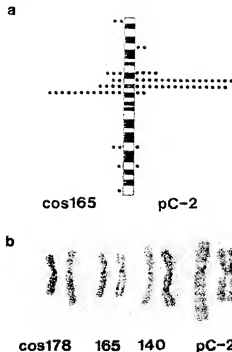


Fig. 5. Localization of the W contigs by *in situ* hybridization. In (a) the grain distribution on chromosome 2 is shown; on the left side for the W-derived cosmid clone cos165 and on the right side for the C₂ probe pC-2. For the probes cos178, cos140 and m654-1 the same localization was found as shown for cos165. In (b) arranged pairs of chromosome 2 are shown after hybridization with the indicated probes. Cosmid probes were detected by a non-radioactive procedure, pC-2 and m654-1 after radioactive labelling.

amplicons I–III, which show that of 29 sites mapped within a stretch of 40 kb only three sites differ between amplicons II and III. It is very likely that the three differences were created by a single event only, which led to a deletion of 1 kb in amplicon II (Figure 1). More differences are found between amplicons II and I (10 out of 33 sites differ) and between III and I (seven out of 33 sites differ). The evolutionary relationship of amplicon IV to the other three amplicons is hard to prove with this strategy, as those parts of amplicons II and III that are homologous to amplicon IV have not yet been cloned (Figure 1).

A chain of events which could have led to the generation of the W regions during evolution is shown in Figure 6. One assumption of the scheme is that the W region precursor had a structure similar to that part of Wb which contains the genes W5 to W9. This is reasonable to assume, as the five genes have the same transcriptional orientation (Figure 1), which also holds true for most genes of the α locus, from where the W precursor must be derived. In a first step, a section of ~150 kb with the genes W7, W8 and W9 is duplicated in such a way that a large palindrome is formed. The newly generated genes 7', 8' and 9' represent the genes W1, W2 and W3 of Wa. In a second step, which led to the duplication of a large block of DNA of ~300 kb, a large palindrome is again formed. A third copy of the genes 7, 8 and 9 (7'', 8'' and 9'') is generated; these copies represent W4, W10 and W11 of Wb. Amplicon IV is a copy of the 3' part of amplicon I, which has been formed in the first step.

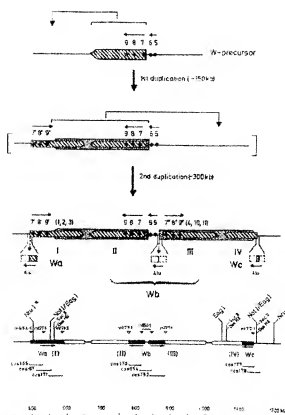


Fig. 6. Model of the generation of the W regions in a multistep process. The precursor contained five genes, marked by filled dots. Arrows indicate transcriptional orientations of genes. The section which becomes duplicated in the course of the first amplification event and the generated amplicons are drawn as hatched arrows with the arrow-heads pointing to the 3' end with respect to the orientation of the genes. The brackets indicate that this structure does not exist in the genome anymore. That part of the intermediate product which is duplicated in a second event and the resulting amplicons are marked by dotted arrows. The four generated amplicons are termed I–IV in accordance with Figures 1 and 3. Genes generated in the first step are marked by ' , those of the second step by ' '. The gene numbers of the present day W regions are given in parentheses. Parts of Alu repeats that have been detected at the joints are indicated. At the amplicon I joint, the 5' part of an Alu repeat is part of the amplicon (hatched rectangle), the deleted 3' part is marked by an open rectangle with a dotted line. At the joint of amplicon III, the 3' part of an Alu repeat is present (open rectangle); it is part of the non-amplified DNA 5' of amplicon III. At the amplicon IV joint only a few nucleotides of an Alu repeat are present (Figure 4c). The deleted 5', as well as 3', sequences are marked by open rectangles with dotted lines. The lower panel shows a long-range map of the W regions, similar to that shown in Figure 3. The open parts of the arrows, which indicate amplicons I–IV, represent those amplified DNA stretches that have not yet been cloned.

A remarkable feature of this series of amplifications is that in the first, as well as in the second, step only one additional copy is formed. This is in contrast to the generation of other amplified units, where the first step of selection yields only a few additional copies; second step selections, and, hence, secondary amplifications, however, usually result in high copy numbers (Saito *et al.*, 1989). Alternatively, the low W copy numbers can be explained by secondary deletions of most of the W copies formed in the amplification events.

Transposition and amplification of the W regions

For the generation of the W regions we propose the following chain of events: (i) the W region precursor, containing the genes W5 to W9 (Figure 6), was transposed to the long arm of chromosome 2 and (ii) the new chromosomal location of the precursor somehow promoted its stepwise amplification.

A pericentric inversion, involving chromosomal bands 2q1 and 2p1, could have been responsible for the transposition of the five V_h genes from the α locus on the short arm to the long arm of the chromosome. Such a chromosomal rearrangement is observed at the evolutionary progenitor of the chimpanzee (Yunis and Prakash, 1982). Transposition of V_h genes by such a process is consistent with a finding by Graninger *et al.* (1988), who detected another part of the α locus, a copy of the α -deleting element (α de); Simionovitch *et al.* (1985), on the long arm of chromosome 2 in 2q11 (Graninger *et al.*, 1988). As 2q11 is the same band in which the W regions are located, one can speculate that the two regions, the W precursor and α de, were transposed to the long arm by the same process. Whereas a copy of the α de is still located 23 kb 3' of C_h (Klobeck and Zachau, 1986), we do not have any evidence that a copy of the W regions exists within the present day α locus.

As an alternative to the transposition by a pericentric inversion, the V_h genes could have been transposed via episomes. Such a mechanism has been proposed to be responsible for the amplification of DHFR and mdrl amplicons in some cell lines (Carroll *et al.*, 1988; Ruiz and Wahl, 1988). While an episome mediated mechanism seems likely for the orphan V_h genes on chromosomes 1, 22 and other chromosomes, we prefer for the W regions the idea of a pericentric inversion.

Data supporting the second assumption, i.e. the new chromosomal location promoted gene amplification, have been reported by Wahl *et al.* (1984), who found an influence of the chromosomal position of transfected CAD genes on the frequency of CAD amplification.

For the amplification event itself, resulting in the formation of palindromic structures, mechanisms such as those recently reviewed by Stark *et al.* (1989) may be responsible. However, to explain the generation of only one additional copy per amplification step, one has to postulate that no replication processes yielding high copy numbers took place in the course of the W region amplification.

Concluding remarks

Pericentric inversions of chromosome 2 involving the chromosomal segments 2p11–2q13 are observed by cytogenetic methods in about 0.1% of today's population (Djalili *et al.*, 1986 and earlier literature). Also, *de novo* inversions have been observed at chromosome 2 (Vejslev and Friedreich, 1984), indicating that the inversion is a spontaneous event. We consider the possibility that extensive sequence homologies between parts of the α locus on 2p12 and the α locus derived W and α de regions on 2qcen–q11 promote the inversions. Interstitial telomere-like repeats, found in 2q11–2q14 (Allshire *et al.*, 1988), might also contribute to the spontaneously occurring pericentric inversions observed in the present day human population.

One of the structural features of the W regions, i.e. the organization of different copies as large palindromes, is also found for the α locus itself (Lorenz *et al.*, 1987). It is tempting to speculate that a duplication involving large parts

of the α locus followed the same molecular mechanisms as the amplification of the transposed W precursor. Proving this hypothesis should be feasible by identifying and analysing the junctions between duplicated and non-duplicated sections, and by cloning the head-to-head junction of the two copies forming the α locus.

Materials and methods

Recombinant DNA and restriction maps

The recombinant cosmids were from libraries described by Pohlenz *et al.* (1987) and Lorenz (1989). Colony hybridization was performed as described previously (Klobeck *et al.*, 1987). Restriction maps and subclones were constructed using established methods (Maniatis *et al.*, 1982). Cosmid clones were characterized as described in Pohlenz *et al.* (1987).

Nucleic acid hybridization and copy number determination

DNA transfer was performed according to Reed and Mann (1985), except for PFG, where the protocol of Rigaud *et al.* (1987) was used. Final washing of filters after hybridization was at 68°C with 40 mM phosphate, pH 7.2, 1% sodium dodecylsulphate.

For copy number determinations of amplicons, the insert of clone m54-1/14 (Figure 2a) was isolated as a 2.2 kb *EcoRI*–*BamHI* fragment on an agarose gel and labelled according to Feinberg and Vogelstein (1983). Digested genomic DNA (10 µg) was electrophoresed, transferred to filters and hybridized. After exposure for 1 day, bands were cut out. For blank values, filter strips above and beneath each band were used. The filter bound activities were measured in a liquid scintillation counter.

PFG

Long-range mapping of DNA from the prostate carcinoma cell line PC-3 (Kaighn *et al.*, 1979) using rare cutting enzymes and orthogonal field gel electrophoresis was performed as described previously (Lorenz *et al.*, 1987).

DNA sequencing

For sequencing of M13-subclones the 'Sequenase' DNA sequencing kit (US Biochemical Corp., Cleveland, OH) was used according to the manufacturer's instructions.

Chromosome bending and in situ hybridization

Metaphase chromosomes were prepared from PHA-stimulated blood lymphocytes. The probes m54-1 and pC-2 were radiolabelled with [³²P]dATP and [³²P]dCTP, and hybridization was done as described in Adolph *et al.* (1987). A non-radioactive detection method was used for the localization of cosmid probes. The cosmid clones were linearized and labelled by the random priming technique with biotinylated dUTP-11. Repetitive DNA sequences within the genomic insert were saturated by prehybridization with a 100-fold excess of human Cot-1 DNA for 4 h (Landegren *et al.*, 1987). The normal *in situ* hybridization was performed after prehybridization.

For probe detection the slides must remain humid. They were rinsed in BT buffer (0.1 M sodium bicarbonate, 0.1% Triton-X-100 pH 8.0) and preincubated for 5 min in BT, 0.1% non-fat dry milk. The slides were treated with peroxidase labelled streptavidin (Enzo Biochemicals Inc., New York; 20 µl per 450 µl BT) and probe detection was done with diaminobenzidine according to the protocol supplied by the manufacturer. The slides were counter-stained with methylene green.

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Human immunoglobulin $C_{\lambda}6$ gene encodes the Kern⁺Oz⁻ λ chain and $C_{\lambda}4$ and $C_{\lambda}5$ are pseudogenes

(isotypes/allotypes/Bence Jones proteins/constant region genes)

P. DARIAVACH, G. LEFRANC, AND M.-P. LEFRANC

Laboratoire d'Immunogénétique, Centre National de la Recherche Scientifique Unité Associée 1191, Génétique Moléculaire, Université des Sciences et Techniques du Languedoc, Place E. Bataillon, 34060 Montpellier Cedex, France

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ABSTRACT Six nonallelic immunoglobulin λ constant region genes have been previously characterized on a 40-kilobase stretch of DNA. The nucleotide sequences of the three upstream genes of this cluster ($C_{\lambda}1$, $C_{\lambda}2$, $C_{\lambda}3$) have been determined by other workers and shown to encode, respectively, the isotypic Mcg, Kern⁺Oz⁻, and Kern⁺Oz⁺ constant region of the λ chains. In this paper, we report the sequence of the three downstream genes of this cluster and show that two of them ($C_{\lambda}4$ and $C_{\lambda}5$) are pseudogenes. However, $C_{\lambda}6$ encodes a Kern⁺Oz⁻ chain and corresponds to the fourth isotype described among the λ proteins sequenced so far. A potentially active J_{λ} (joining) segment, with the canonical heptamer and nonamer sequences for rearrangement, is located 1.5 kilobases upstream of $C_{\lambda}6$. The amino acid sequence encoded by the $C_{\lambda}6$ gene is compared with the constant region sequences of various monoclonal Bence Jones λ proteins. Allotypic and isotypic differences confirm the polymorphism and complexity of the human C_{λ} locus.

In humans, the constant (C) region of the immunoglobulin λ light chains consists of at least four nonallelic or isotypic forms that differ by limited amino acid substitutions to produce the serological markers Kern (Ke) (1, 2), Oz (3-5), and Mcg (6, 7). Several additional substitutions have been described (8-16), but it is unknown whether these represent allelic variants or distinct isotypes. The human immunoglobulin λ light chain genes have been mapped to chromosome 22 (17) at band q11 (18, 19), and six nonallelic C region genes ($C_{\lambda}1$ to $C_{\lambda}6$) have been characterized on a 40-kilobase (kb) stretch of DNA (20). The number of C_{λ} genes varies between six and nine per haploid genome (21). These variations were detected by restriction fragment length polymorphism (21) and seem to have arisen from unequal meiotic crossing-over with a duplication of the $C_{\lambda}2$ and $C_{\lambda}3$ genes. Moreover, three additional C_{λ} -like genes have been recently identified, which map on different stretches of DNA and are nonallelic (22). One of these is a pseudogene, whereas the two others encode a putative λ chain C region whose sequence differs from that of the λ chains described so far.

Only three C_{λ} genes ($C_{\lambda}1$, $C_{\lambda}2$, and $C_{\lambda}3$) belonging to the cluster described by Hieter have been sequenced (20), and they have been shown to encode, respectively, the Mcg, Ke⁺Oz⁻, and Ke⁺Oz⁺ C region of the λ chains. In this paper, we report the sequences* of the three genes located downstream in this cluster and show that two of them ($C_{\lambda}4$ and $C_{\lambda}5$) are pseudogenes, whereas $C_{\lambda}6$ encodes a Ke⁺Oz⁻ chain, the fourth isotype described among the proteins sequenced so far. This $C_{\lambda}6$ gene has a potentially active J_{λ} joining region, with the canonical heptamer and nonamer sequences for rearrangement, 1.5 kb upstream of the coding C region.

MATERIALS AND METHODS

Construction of a Phage Library from LY67 DNA. DNA prepared from LY67 cells (a λ -producing Burkitt's lymphoma) (23) was partially digested with *Mbo* I. Restriction fragments 15-20 kb long were ligated into *Bam*HI-digested DNA of phage λ 2001 (24) and packaged *in vitro*. Recombinant phages were screened by the *in situ* plaque hybridization procedure (25).

Probes. A genomic clone (Chr 22A5) in λ gt-10WES (26) was kindly provided by T. H. Rabbitts (Medical Research Council, Cambridge, England). This clone contains an 8.0-kb *Eco*RI fragment that includes the known nonallelic Ke⁺Oz⁻ ($C_{\lambda}2$) and Ke⁺Oz⁺ ($C_{\lambda}3$) genes and the flanking sequences (20). We subcloned a 700-base-pair (bp) *Bgl* II-*Eco*RI fragment containing only the Ke⁺Oz⁻ $C_{\lambda}3$ gene (Fig. 1), and this $C_{\lambda}3$ probe cross-hybridizes with all the other C_{λ} -like genes (20). It was radioactively labeled with [α -³²P]dCTP by nick-translation (27) and was used to screen the LY67 phage library.

Subcloning and Sequencing Strategies. One clone, LY67 $C_{\lambda}3$ -6 (Fig. 1), was shown to contain $C_{\lambda}3$ to $C_{\lambda}6$. Appropriate subclones were made in pUC vectors (29). Nucleotide sequence analysis was carried out by dideoxy chain-termination procedures (30) in M13 vectors (31) by deploying exonuclease III-nuclease S1 methods (32) or directed sequencing using known restriction enzyme sites.

Oligonucleotide Synthesis and Hybridization. A 19-mer oligonucleotide 5' GTGTTCCGGCGAGGGACCA 3' corresponding to part of the $J_{\lambda}3$ gene segment sequence (this paper and ref. 28) was synthesized, radiolabeled, and hybridized to the LY67 $C_{\lambda}3$ -6 clone to search for other J_{λ} segments. Low-stringency washes were carried out at room temperature.

RESULTS

Rearrangement of a $V_{\lambda}III$ Subgroup Gene to $J_{\lambda}3$ in the LY67 Cell Line. One clone (LY67 $C_{\lambda}3$ -6) containing a 18-kb piece of genomic DNA was isolated and characterized. A restriction map of this clone is shown in Fig. 1. Comparison of this map with one previously published (20) suggested that this clone contains four C_{λ} genes, namely $C_{\lambda}3$ to $C_{\lambda}6$. The sequence of the 5' end of the LY67 $C_{\lambda}3$ -6 clone shows that a V_{λ} gene rearrangement has occurred, joining this gene to the $J_{\lambda}3$ gene segment, which is located 1.5 kb upstream of $C_{\lambda}3$ (28). Fig. 2A shows the partial nucleotide sequence of the rearranged V_{λ} in LY67 and that of a V_{λ} gene assigned to the $V_{\lambda}III$ subgroup and isolated from the Burkitt lymphoma cell line PA682 (28).

Abbreviations: C, constant; J, joining; V, variable; Ke, Kern.
*The sequences reported in this paper are being deposited in the EMBL/GenBank data base (Bolt, Beranek, and Newman Laboratories, Cambridge, MA, and Eur. Mol. Biol. Lab., Heidelberg) [accession nos. J03009 ($C_{\lambda}4$), J03010 ($C_{\lambda}5$), and J03011 ($C_{\lambda}6$)].

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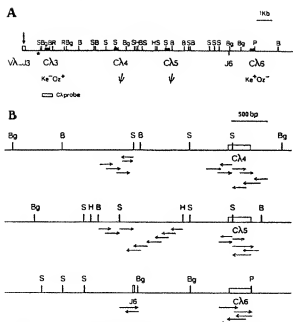


Fig. 1. (A) Restriction map of LY67 C3-6 clone. (B) Sequencing strategy. B, BamHI; Bg, Bgl II; S, HindIII; R, EcoRI; S, Sst I. Of the Pst I sites (P), only the one used for subcloning the fragment containing the C₆ gene is indicated. The rearrangement V-J₃ is indicated by an arrow (V, variable region). An asterisk shows the location of a polymorphic BamHI site present in PA682 DNA (28) but absent from our LY67 clone.

The deduced amino acid sequence of the rearranged V gene of LY67 is also compared to the V region of the protein DEL of the subgroup V_hIII (33, 34). A 75% sequence identity indicates that the V_h gene rearranged in LY67 is a member of the V_hIII subgroup gene family, and this is in agreement with the detection of a transcript hybridizing to a V_hIII probe in the

LY67 cell line (28). The J₃ segment of the LY67 C3-6 clone, compared to the J₃ segment rearranged in PA682, shows two nucleotide differences (one of them resulting in a valine/leucine amino acid substitution) that may be due to allelic polymorphism. Two other nucleotide differences are observed at the V-J junction and are probably explained by a flexibility in the mechanism by which junctions occur (35, 36).

C₆ Encodes a Ke⁺Oz⁻ Chain. Fig. 2B shows the nucleotide sequence of the C₆ gene and the encoded amino acid sequence (106 residues). The residues Ala, Ser, and Thr, found, respectively, at codons 6, 8, and 57 (positions 112, 114, and 163 according to ref. 34) indicate that C₆ encodes a Mcg⁺ protein. Arg (codon 83, position 190) corresponds to the Oz⁻ marker, whereas Gly (codon 46, position 152) characterizes the Ke⁺ marker. Therefore the C₆ gene encodes the fourth isotype Ke⁺Oz⁻.

J₆ Segment Is 1.5 kb Upstream of C₆. Only the J₆ (22) and J₃ segments (ref. 28 and this paper) have been characterized; they have been localized in genomic DNAs at 1.5 kb upstream of the respective C₄ coding regions. We therefore used an oligonucleotide corresponding to the J₃ sequence (see Materials and Methods) to search for homologous J₆ segments in the LY67 C3-6 clone. As expected, a strong signal was obtained for the J₃-containing fragments, whereas a weaker signal allowed us to detect the J₆ segment in a Sac I-Bgl II fragment upstream of C₆. The sequence of this J₆ segment (Fig. 2C) showed that it encodes 12 amino acids (among them the characteristic Phe-Gly-Xaa-Gly residues) and that it also possesses the canonical heptamer and nonamer sequences essential to V-J rearrangement (37, 38). No signal corresponding to the putative J₄ and J₅ segments could be detected in the LY67 C3-6 clone by using either the oligonucleotides (J₃ probe) or the genomic Sac I-Bgl II fragment (J₆ probe), indicating that if these segments exist their homology is too weak to be detected in our conditions of hybridization. Since the J₃ and J₆ gene segments are 1.5 kb upstream of their respective coding regions, we subcloned fragments located, respectively, at about the same distance upstream of C₄ and C₅. Although in both cases, we detected some conserved heptamer sequences, we did not

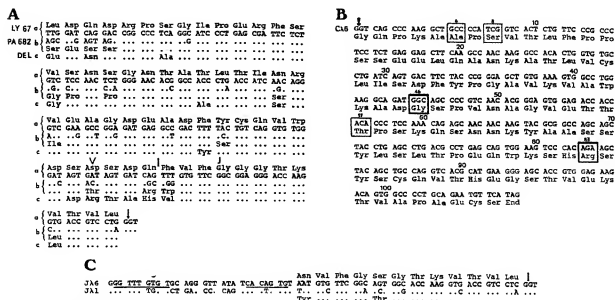


Fig. 2. Nucleotide and amino acid sequences. (A) Partial sequence of the LY67 V-J₃ rearranged gene. (B) Sequence of the C₆ gene. (C) Sequence alignment of J₆ and J₁ (22).

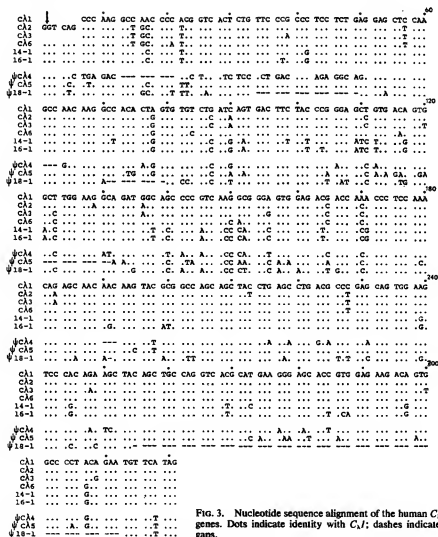


Fig. 3. Nucleotide sequence alignment of the human C_1 genes. Dots indicate identity with C_1 ; dashes indicate gaps.

find the characteristic Phe-Gly-Xaa-Gly residues or the expected splice site at a downstream position. It is possible that the heptamer sequences are attached to poorly conserved pseudo J_4 segments and we cannot exclude the possibility that the putative J_4 and J_5 are localized in

fragments that were not sequenced, upstream of the C_4 and C_5 genes.

C_4 and C_5 Are Pseudogenes. Nucleotide sequences and the encoded amino acid sequences of C_4 and C_5 are shown in Figs. 3 and 4. Both genes are pseudogenes; the third codon of C_4 is a stop codon, and C_5 displays three deletions. The first deletion of 9 bp spans codons 5 to 7 and the other two deletions excise codons 21 and 64. C_5 has an 11-bp deletion (codons 41–44) resulting in a frameshift.



Fig. 4. Protein sequences derived from the human C_1 gene sequences. The standard one-letter symbols are used.

Table 1. Amino acid differences between the four nonallelic forms of the human C_1 regions

Isotype	Gene	Amino acid residue			
		112 (6)	114 (8)	152 (46)	190 (83)
Mg ²⁺ Ke ⁺ O ₂ ⁻	C_1	Asn	Thr	Gly	Lys
Mg ²⁺ Ke ⁺ O ₂ ⁻	C_2	Ala	Ser	Ser	Thr
Mg ²⁺ Ke ⁺ O ₂ ⁻	C_3	Ala	Ser	Ser	Thr
Mg ²⁺ Ke ⁺ O ₂ ⁻	C_4	Ala	Ser	Gly	Thr

Residue numbering is according to ref. 34; parentheses enclose numbering of the codons in the C_1 genes.

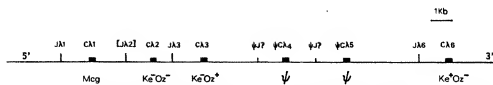


FIG. 5. Physical map of the human λ light chain C region. $C_{\lambda 1}$, $C_{\lambda 2}$, and $C_{\lambda 3}$ correspond, respectively, to the nonallelic Mcg, $K\epsilon^*Oz^-$, and $K\epsilon^*Oz^+$ chains (ref. 20; see Table 1). $C_{\lambda 4}$ and $C_{\lambda 5}$ are pseudogenes (ψ), whereas $C_{\lambda 6}$ encodes a $K\epsilon^*Oz^+$ chain. $J_{\lambda 1}$ (22), $J_{\lambda 3}$ (ref. 28 and this paper), and $J_{\lambda 6}$ (this paper) have been localized 1.5 kb upstream of $C_{\lambda 1}$, $C_{\lambda 3}$, and $C_{\lambda 6}$, respectively. $J_{\lambda 2}$ has not yet been localized in genomic DNA. No $J_{\lambda 1}$ gene segment has so far been identified upstream of $C_{\lambda 4}$ and $C_{\lambda 5}$.

DISCUSSION

In human λ chain C regions, the Mcg marker involves amino acid residues at positions 112, 114, and 163 (numbering according to ref. 34; Table 1) corresponding, respectively, to codons 6, 8, and 57 of the C_{λ} genes. Mcg⁺ proteins have residues Asn-112, Thr-114, and Lys-163, whereas Mcg⁻ proteins have residues Ala, Ser, and Thr, respectively, at these locations. However, the recently sequenced Mor protein is different from the other Mcg⁺ proteins by having Ala-163 instead of Thr-163 (16). The Ke and Oz markers occur at positions 152 and 190, respectively: $K\epsilon^*Oz^-$ proteins have Gly-152 and $K\epsilon^*Oz^+$ have Ser-152; Oz⁻ proteins have Lys-190 and Oz⁺ Arg-190. These markers define four nonallelic forms of human λ chain C regions (Mcg, $K\epsilon^*Oz^-$, $K\epsilon^*Oz^+$, and $K\epsilon^*Oz^+$), which are encoded, respectively, by the $C_{\lambda 1}$, $C_{\lambda 2}$, $C_{\lambda 3}$ (20), and $C_{\lambda 6}$ (this paper) genes (Fig. 5).

In Fig. 6, monoclonal Bence Jones proteins have been assigned as products of the C_{λ} genes 1, 2, 3, or 6 on the basis of the presence or absence of residues characteristic for the Mcg, Ke, and Oz markers. In most cases, there is complete concordance between the protein and the deduced amino acid sequence of the corresponding C_{λ} gene. However, other amino acid changes have been found in several proteins (Table 2 and Fig. 6). Since these substitutions have been noted only once, they could represent allelotypic differences. However, it is not excluded that some of the proteins $K\epsilon^*Oz^-$ could be encoded by a C_{λ} gene resulting from the duplication of the $C_{\lambda 2}$ - $C_{\lambda 3}$ region, as has been described in some individuals (21). In such cases these sequences should represent new isotypic differences due to the presence of several nonallelic copies of $C_{\lambda 2}$ gene. Differences observed

Table 2. Sequence variations in the C region of human λ chains

Isotype	Common amino acid	Residue number	Variant amino acid	Protein	Ref.
Mcg	Lys	156	Glu	WEIR	8
$K\epsilon^*Oz^-$	Lys	129	Glx	CH	9
			Arg	NIG68	10
			Ser	ATK	11
	Ala	143	Val	MZ	12, 13
	Val	155	Ile	HIL	14
	Lys	156	Glu	SA	11
	Lys	172	Asn	MZ	12
$K\epsilon^*Oz^+$	Lys	187	Gln	ATK	11
	Gln	195	Leu	NIG68	10
$K\epsilon^*Oz^+$	Asp	151	Glu	EV	15

Residue numbers according to Kabat *et al.* (34).

in the $J_{\lambda 2}$ sequences (Fig. 6) might for the same reason be either allelotypic or isotypic. Differences in the $J_{\lambda 1}$ segment region might represent allelotypic differences, although the presence of not yet identified other $J_{\lambda 1}$ segments cannot be ruled out.

If we compare the deduced amino acid sequence of $C_{\lambda 6}$ gene with two known $K\epsilon^*Oz^+$ proteins that have identical C_{λ} coding regions, SM (53) and Kern (52), the protein predicted for $C_{\lambda 6}$ shows three differences: (i) lysine at position 145 (codon 39) instead of threonine, (ii) asparagine at position 152 (codon 50) instead of lysine, and (iii) alanine at position 212

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FIG. 6. Sequences of C regions of human λ chains. The protein sequences encoded by the four "active" C_{λ} genes and associated J_{λ} gene segments are compared with the λ protein C regions. The numbering of the C_{λ} region amino acids is according to ref. 34—e.g., positions 169, 201, and 202 are excluded in the C_{λ} sequences for purposes of alignment with human C_{λ} chains. For an easier alignment, the J_{λ} and C_{λ} gene segments are considered as being spliced and a vertical line is drawn to indicate the V-J junction. The horizontal line in the SM sequence shows a deletion.

(codon 103) instead of threonine (Fig. 6). These differences may represent allelotypic variations, although we cannot entirely exclude the possibility that these different Ke^oOz⁺ sequences are encoded by nonallelic genes. More sequences of C₁ genes or λ proteins should help estimate the extent of the human λ chain polymorphism.

Note Added in Proof. The J₂ gene segment has recently been localized upstream of C₂ (54).

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Evolutionary and structural influences on light chain constant (C_L) region of human and mouse immunoglobulins

(structure-function correlation and evolution/ λ - κ comparisons/comparative structures)

ELVIN A. KABAT*, EDUARDO A. PADLAN†, AND DAVID R. DAVIES†

* Fogarty International Center and † Laboratory of Molecular Biology, National Institute of Arthritis, Metabolism and Digestive Diseases, National Institutes of Health, Bethesda, Maryland 20814; and ‡ Departments of Microbiology, Human Genetics and Development and Neurology, Columbia University and the Neurological Institute, Presbyterian Hospital, New York, N.Y. 10032

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ABSTRACT A comparison of five constant region sequences of human and mouse κ and λ immunoglobulin chains has been undertaken in order to reveal sequence homologies and evolutionary relationships. Simultaneously, a comparison with the three-dimensional structure of one mouse κ -chain (McPC 603) has suggested structural reasons why many of the residues are invariant or conserved along κ versus λ lines. There are a number of residues that have remained invariant despite exposed positions for reasons that do not appear to be connected with the folding of this C_L domain.

The constant region (C_L domain) of immunoglobulin light (L) chains contributes substantially to the functioning of an immunoglobulin molecule. While it is not involved directly in the specificity and complementarity of the antibody combining sites, it is joined to its counterpart in the heavy (H) chain, the C_{H1} domain, by a variety of noncovalent interactions as well as in most immunoglobulins by a disulfide bond at its C-terminal or subterminal Cys. This -S-S- bond is not essential to L-H association, which is maintained noncovalently even after reduction and alkylation. In one immunoglobulin subclass of IgA2, the L-H bond does not occur, but an L-L dimer is formed, which remains noncovalently linked to the H chains. Bence Jones proteins in the form of L-L dimers also occur and may be held together by noncovalent forces (1).

There are two subclasses of light chains, κ and λ , which are present in almost all species examined; both are found in all five classes of immunoglobulins, IgG, IgM, IgA, IgD, and IgE, but each immunoglobulin molecule contains two identical κ or two identical λ chains.

The availability of the sequences of the C_L domain of human κ , human λ , mouse κ , and two mouse λ light chains (2), together with x-ray data on the three-dimensional structure of this domain (3-5) made it desirable to evaluate, if possible, structural influences on the evolution of these domains.

The present study attempts, residue by residue among these five chains, to relate preservation or variation of sequence to structure and function as evaluated from a three-dimensional model of the C_L and its interactions with the C_{H1} domain. The findings show some interesting stretches of sequence in which invariance predominates, and others in which evolutionary divergence has been essentially along κ

and λ lines. Positions at the surface of protein molecules that are accessible to solvent may undergo many mutational changes which do not affect three-dimensional folding, while residues that are buried tend to be invariant or highly conserved (6). In C_L domains mutations involving residues that contact the C_{H1} domain also tend to be restricted.

The present study shows that residues preserved along κ and λ lines generally show conservative substitutions if in the interior of the domain or if buried. Fewer exposed residues which have diverged along κ and λ lines are homologous. Certain residues remain invariant despite an essentially exposed position and for no obvious reason. Of special interest is the observation that at only two and four positions, respectively, were human κ identical with human λ and mouse κ identical with mouse λ , while human κ and mouse κ were identical at 29 and human λ and mouse λ at 39 positions.

MATERIALS AND METHODS

The model of the Fab fragment of mouse McPC 603 constructed from x-ray data at 3.1 Å resolution was used (5) as well as published information on the C_L regions of a Bence Jones dimer (4) and human Fab fragment (3). Sequences of human κ , human λ , mouse κ , and two mouse λ chains were available (2). These sequences were aligned for maximum structural and sequence homology from residues 101 to 215 and modified to include the additional data reported (7). Each residue was located in the model of McPC 603 and classified according to whether it was: exposed, 0; mainly exposed, 1; partly exposed, partly buried, 2; mainly buried, 3; completely buried, 4; or in contact with C_{H1} , C. In addition, each position was classified as: invariant; four of five chains and three of five chains identical; human κ and human λ identical; mouse κ and mouse λ identical; human and mouse κ identical; human and mouse λ identical; and human κ , human λ , mouse κ and mouse λ different.

RESULTS

Table 1 lists the sequences of the five chains from positions 101 to 215. Above each residue is its classification from its position in the model.

Fig. 1 summarizes the sequence data in Table 1 with respect to the identities specified above. It is evident that clusters of invariant residues and those with 4/5 chains identical occur, notably at positions 118 to 123, 148 to 152 (excluding 150), 176 to 182 (180 3/5 identical), and 194 to 200 (exclud-

Abbreviations: C and V, constant and variable regions of immunoglobulin chains; L and H, light and heavy chains of immunoglobulins.

Table 2. Location from x-ray structure and degree of evolutionary preservation of residues in the C₁ domain and switch region of human and mouse light chains

Location	Code	Invariant	4/5	3/5	Identities—number of residues				
					Human κ Human λ	Mouse κ Mouse λ	Human κ Mouse κ	Human λ Mouse λ	Human κ , Human λ Mouse κ , Mouse λ different
Exposed	0	7	7	3	1	2	10	16	6
Mainly ex- posed	1	2	2			1	4	4	2
Partly ex- posed, partly buried	2	1	4	1			2	3	
Mainly buried	3	2		2			2	1	
Completely buried	4	11	4	1			3	5	
Contacts H chain	C	5		1	1	1	8	7	
Total		28	17	8	2	4	29	36 [†]	8

[†] Excludes residues with * and • in Fig. 1 and residue 215 not specifiable; gaps in κ at position 108 and in λ chains at positions 169 and 201 and 202.

these included the Inv marker 191. Position 152, which in human λ chains carries the Kern marker, was also exposed (3).

Of the 28 invariant residues 18 were contact residues or were mainly or completely buried, while nine were mainly or completely exposed. Considering together the 25 positions in which 4/5 and 3/5 residues were identical, 12 were completely or mainly exposed, only one was a contact residue, while five were completely buried.

Of the 29 positions at which human κ and mouse κ had the same amino acid and the six (footnote Table 2) at which human λ and one of the mouse λ chains had the same amino acid, seven in each were contacting residues and three and six, respectively, were completely buried while 15 and 20, respectively, were completely or mainly exposed.

Of the seven invariant exposed residues six were charged, 2 Glu, 1 Asp, and 3 Lys; the seventh was Cys 214 which forms the S-S bond to the H-chain or in some cases to another L-chain. The location of these six residues and the two mainly exposed residues His and Thr in the model provides no insight into the basis for their invariance.

The remaining 19 invariant residues, those partly or completely buried and the contacting residues, included 12 strongly hydrophobic residues, 1 Trp, 2 Phe, 4 Pro, 3 Leu, and 2 Val; the remaining residues were Cys 134 and Cys 194, which form the domain S-S bond, two less strongly hydrophobic residues Ala and Thr, and three Ser.

Examining the residues with 4/5 and 3/5 identities, the partly, completely buried, and contacting residues were also overwhelmingly hydrophobic, consisting of 2 Tyr, 5 Val, 1 Thr, and 1 Ala, the others being 1 Gly and 1 His and the gap at position 201. The nonidentical residues at these positions were also hydrophobic in almost all cases, involving replacements of 2 Ile, 2 Leu, and Ala for the five Val residues and of Phe for one Tyr; the remaining Tyr 192 was replaced by Ser in both mouse λ chains, the Thr by Ser or His, the Ala by

Asx, and the Gly by Thr. There is obviously an extraordinary preservation of structure in terms of these groups of residues.

Among the 15 and 16 residues that have diverged along κ versus λ lines and which are partly, mainly, and completely buried or are contacting residues, eight are identical pairs, three of these involve conservative hydrophobic substitutions, Ile-Leu, Val-Leu, and Leu-Ile, at positions 117, 132, and 136, respectively, involving mainly and completely buried residues; the remaining pairs are the substitutions Thr-Lys at 172, Glu-Glu 124, Ser-Thr 131, Glu-Ser 165, and Thr-Tyr 178, the last four being contacting residues. The presence in κ chains of an additional residue at 169 causes Thr 172 to be completely buried in the mouse C₁ domain; in the λ chains, Lys 172 is probably completely exposed to solvent. The unpaired residues are 1 Phe, 1 Tyr, 2 Ser, 1 Asp, and 1 Thr in the κ , and 1 Ala, 1 Val, 1 Glu, 1 Glu, 1 Lys, and 3 Thr in the λ group.

The κ versus λ differences show a predominance of completely and mainly exposed residues with 14 of 29 and 20 of 36 residues in κ and λ , respectively, falling into this group; nine are pairs.

The region 101–108 consists of two invariant residues, 101 and 102, followed by six positions in which residues with 4/5 identities alternate with residues which have evolved along κ and λ lines, including the gap at position 108; four of the positions are completely exposed, one is partly exposed, and two are completely buried. Arg 107 marks the end of the mouse V₁ domain; C₁ starts with Ala 109. The additional residue at position 108 in λ chains could be accommodated by a hairpin bend facilitated by Gly 107 or by Pro 109.

There is a cluster of invariant residues from 118 to 123 consisting of three contacting residues, 118, 119, and 121, one partly buried 120, and two exposed residues, 121 Ser (4/5) with an Asp alternative, and an invariant Glu 123. The region 124–140 contains some invariants but is largely made

up of residues which have evolved along κ and λ lines; residues 130-137 consist exclusively of contacting and completely buried residues of which three are invariant, in another three κ and λ differ, and in the others more variation has occurred. At 135 human κ and λ have Leu while mouse κ has Phe and mouse λ Thr and at position 137 human and mouse κ have Asn, human λ Ser, and mouse λ Thr.

The most striking region which has been preserved along κ versus λ lines, 160 to 175, has five contacting, one completely and four partly buried residues, and two mainly and three completely exposed residues. It is followed by a largely invariant cluster, 176 to 181, of contacting and buried residues. The remainder of the molecule is mainly exposed except for the region around Cys 194, in which buried and mainly exposed residues alternate and there is no clustering of invariant or κ versus λ residues.

DISCUSSION

The general structural relationships described for the C_L region of human and mouse κ and λ chains are an initial attempt to understand the basis for the evolutionary preservation of certain regions as essentially invariant and the preservation of others along κ versus λ lines, an evolutionary divergence which took place about 200 million years ago (8). The principles established from sequence and structural findings on other proteins generally apply equally well to the immunoglobulins. Thus the buried residues and those contacting the heavy chain tend to be largely invariant and hydrophobic, while residues that are exposed or mainly exposed may vary, and are generally polar. These also include the few residues that have evolved along human versus mouse lines. There are, however, substantial numbers of hydrophobic residues that are invariant or identical in 4/5 or 3/5 chains and that occur in regions of the molecule for which no obvious structural basis for their conservation may be assigned.

The marked clustering of residues which have been preserved along κ versus λ lines in certain portions of the chain, most notably at positions 160 to 175, or have been maintained invariant, such as 118 to 123, suggests that these may have unique functions.

The C_L domain has been extraordinarily preserved once κ versus λ diversification occurred, since at only eight posi-

tions were all four chains different and at only five other positions were human and mouse chains identical despite a κ versus λ difference (Fig. 1).

The immunoglobulin findings were compared with values for human and mouse hemoglobins, using the α chain as equivalent to κ and the β (γ) chain equivalent to λ (8). Since hemoglobin chains are longer, the data were normalized to 115 residues. The findings for hemoglobin are strikingly different from the immunoglobulin results. Thirty-six residues were invariant, and 4/5 and 3/5 chains were identical at 12 and at 49 positions, respectively. Human and mouse α and human and mouse β were identical at 13 and 11 positions, respectively. At only one position each were mouse α and mouse β , human α and mouse β , and human β and mouse α identical. There were no positions at which all four chains differ. Thus there were 97 residues in which three or more chains were identical in hemoglobins in contrast to 53 such residues in the immunoglobulins. This comparison indicates a higher degree of evolutionary conservation of residues that did not differentiate along α versus β lines than residues that differentiated along κ versus λ lines. The immunoglobulin C_L domain thus shows a higher degree of evolutionary adaptability than do the hemoglobins.

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Commentary
Progress in programming antibody fragments to crystallize

Allen B. Edmundson ^{a,*}, Carl A.K. Borrebaeck ^b

^a Oklahoma Medical Research Foundation, 825 NE 13th Street, Oklahoma City, OK 73104, USA

^b Department of Immunotechnology, Lund University, P.O. Box 7033, S-220 07 Lund, Sweden

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Abstract

Completion of the X-ray analysis of the human B7-15A2 Fab opened a new vista (Immunotechnology 3, no. 4). In the crystal lattice, both the λ -type light chain (C_L domain) and $\gamma 1$ -type heavy chain (C_H1 domain) participated in formation of antiparallel β -pleated sheets with neighboring molecules related to the reference Fab by 2-fold axes. This observation evoked memories of the first description of this type of packing for human Bence-Jones (λ chain) dimers 20 years ago (Ely K.R. et al. Biochemistry 1978;17:158–167). Reexamination of packing interactions in selected crystal systems revealed that the C domains of λ and $\gamma 1$ chains were structurally amenable to the formation of such cross-molecule β -structures, but κ chain C_L domains were not. In the latter, a single proline residue disrupted the order of β -strand 3–3 in the middle of the surface used in λ and $\gamma 1$ chains for intermolecular interactions with symmetry-related molecules. For the packing of Fv molecules, the V_L domains are structurally well suited for analogous packing interactions through antiparallel 4–1 β -strands in adjacent molecules. Such interactions have been shown to provide the driving force in the crystal packing of a human (Pot) Fv from an IgM- κ cryoglobulin. Together, these observations suggest several avenues through which propensity to crystallize can be programmed into the designs of synthetic human Fabs, Fvs and single-chain antibodies. © 1998 Elsevier Science B.V.

Keywords: Human B7-15A2; β -pleated sheets; Bence-Jones dimers; Fv molecules

1. Introduction

One of the most fertile sources of information about macromolecular interactions can be found amongst protein constituents orderly arranged in crystal lattices. This resource is regularly being

* Corresponding author. Tel.: +1 405 2711673; fax: +1 405 2713980.

upgraded through an ever increasing number of structures solved by X-ray analysis. Symmetry operations in most of the commonly appearing space groups afford exciting opportunities to observe groups of these molecules in a variety of combinations and orientations within the lattice. Taking advantage of such opportunities is made possible by the powerful computer-assisted graphics workstations and sophisticated software that are currently available.

A particularly elegant stacking pattern was observed in crystals of an Fab from a human IgG1- λ antibody (B7-15A2) against tetanus toxoid. The key design element in these crystals is a continuous ribbon of Fabs stabilized through interlocking, antiparallel six-stranded β -pleated sheets between the C_L and C_{H1} domains of symmetry-related molecules. Each Fab is bound to two other molecules, one on each side, and this process is repeated to produce a ribbon of indefinite length. When correlated with information previously accumulated for human Bence-Jones (L chain) dimers and other Fabs, the crystal packing of B7-15A2 suggests that the propensity to crystallize can be programmed into the design of some synthetic Fabs.

2. Bilateral formation of β -pleated sheets across 2-fold axes

A stereo diagram illustrating the self assembly of B7-15A2 Fabs into a trimeric packing unit in the crystal lattice is presented in the top panel of Fig. 1. Main chain atoms of the outer strand (3–3) of the three-chain antiparallel β -pleated sheet in the C_L domain of one Fab (e.g. central molecule in Fig. 1) form hydrogen bonds across a crystallographic 2-fold axis with constituents of β -strand 3–3 in the C_{H1} domain of a symmetry-related Fab (on the right). On the opposite (left) side of the reference molecule, the interactions are repeated in reverse as C_{H1} interacts with the C_L domain in a third Fab.

Bilateral, six-stranded β -sheet structures across crystallographic dyads were first described for a human (Mcg) Bence-Jones dimer

[1]. They were subsequently observed in the X-ray analysis of another (Loc) λ -type Bence-Jones protein [2]. This type of packing is shown in the lower panel of Fig. 1 for three adjacent subunits in the ribbon of Mcg dimers. Models are displayed in the same format and orientation to facilitate comparisons of the B7-15A2 and Mcg ribbons.

3. Formation of β -pleated sheets by C_L or C_{H1} , but not both

A six-stranded β -pleated sheet is also found in crystals of a *heterologous* dimer of two human λ -chains (Mcg and Hud), but the interactions are not bilateral (Luke Guddat and Allen Edmundson, unpublished results). Instead, cross-molecule hydrogen bonding occurs only between C_L domains of the Mcg subunits. Therefore, the principal packing motif is a dimer of the hybrid and not a continuous ribbon. These relationships are illustrated in Fig. 2. A similar pattern is observed in crystals of the Fab New, where the six-stranded β -sheet spans antiparallel C_{H1} domains [3].

4. Cross-molecule β -sheet produced by κ -type V_L domains

Currently, there is only one known example of V domains engaging in this type of packing. In crystals of an Fv (V_L – V_H combination) from a human (Pot) IgM cryoglobulin, each κ -type V_L domain interacts with an antiparallel V_L domain across a 2-fold axis [4]. Unlike the C_L and C_{H1} packing patterns, the N-terminal β -strand 4–1 of V_L is the principal interacting unit in the Pot Fv. As shown in Fig. 3, the key packing step is again the formation of a dimer rather than a ribbon. This type of packing leads to a crystal with a high fractional volume of protein in the unit cell (very advantageous in X-ray analysis). For instance, the protein:solvent ratio in the Pot Fv crystals was 0.64:0.36, compared with 0.50:0.50 in the B7-15A2 Fab and 0.40: 0.60 in the Mcg L chain dimer.

5. Structural features favoring β -sheet formation

If Fv fragments are excluded, bilateral participation in cross-molecule β -sheet structures seems to be primarily dependent on the relative orientations of C_L and C_H1 . Among human Bence–Jones proteins and human Fabs, the components most likely to manifest such favorable properties are C_L domains from λ -type light chains, alone or in combination with $\gamma 1$ heavy chains. In these domains the 3–3 β -strands are often held in accessible and ordered positions by quite regular hydrogen bonding with 3–2 β -strands along their inner surfaces. On their outer surfaces, the backbone oxygen and nitrogen atoms not involved in such intradomain interactions are free to form hydrogen bonds with the corresponding β -strands in symmetry-related molecules.

The portions of the 3–3 β -strands conducive to docking are comparable in human λ and $\gamma 1$ chains. They involve the first six residues following the turn between β -strands 3–2 and 3–3 and have the sequences Gly–Ser–Thr–Val–Glu–Lys in the λ -type L chains (residues 203–208 in computer numbering; no allowances for gaps or insertions) and Ser–Asn–Thr–Lys–Val–Asp (residues 207–212) in the $\gamma 1$ H chains. There are four intermolecular hydrogen bonds formed between the backbone atoms of antiparallel 3–3 β -strands in the Mcg dimer (both L chains), the Mcg \times Hud hybrid (Mcg subunits only), the B7-15A2 Fab (L and H chains) and the Fab New (H chains only). The pattern in Mcg is simple to understand and will be considered first.

6. Cross-molecule hydrogen bonding in crystals of the Mcg dimer

Val 206 is located at the center of the interactions. Hydrogen bonds 1 and 2 are formed between the main chain amide group and carbonyl oxygen of Thr L205 of monomer 1 and the carbonyl oxygen and amide group of Glu 207 of monomer 2. In a reciprocal manner, the amide and carbonyl groups of Thr L205 in monomer 2 interact with the carbonyl and amide groups of

Glu L207 in monomer 1 to form hydrogen bonds 3 and 4. There is a fifth hydrogen bond between the side chain hydroxyl group of Ser 204 in monomer 2 and the ϵ -amino group of Lys 208 monomer 1. Because of local asymmetry in the conformations, only one hydrogen bond is possible because the side chains of Ser 204 in monomer 1 and Lys 208 in monomer 2 are not sufficiently close to interact.

7. Hydrogen bonding across 2-fold axes in the Mcg \times Hud hybrid

It is interesting that only the Mcg monomer participates in the hydrogen bonding of the Mcg \times Hud hybrid, despite the fact that the amino acid sequences of the two λ -chains are identical in the 3–3 β -strands and surrounding regions of the C_L domains. Again, the pattern of four hydrogen bonds in the L205–207 central region is repeated, as shown in Fig. 2.

8. Intermolecular hydrogen bonding between C_L and C_H1 in B7-15A2

There is a slight variation in the B7-15A2 Fab (see Fig. 1, top panel). At one end of β -strand 3–3, the oxygen atom of Gly L203 makes a hydrogen bond with the amide group of Asp H212 (position analogous to position L207). In the central region, the amide nitrogen and carbonyl oxygen of Thr L205 form hydrogen bonds with the oxygen and nitrogen atoms of Lys H210 (position analogous to L205). At the distal end the amide group of Glu L207 hydrogen bonds with Asn H208 (analogous to residue L203).

9. Hydrogen bonding between antiparallel C_H1 domains in Fab New

In Fab New [3] there are four hydrogen bonds involving Lys H210 and Asp H212 of two C_H1 domains (not shown). It is significant that these residues in the 3–3 β -strands of both New and B7-15A2 H chains are separated from the inter-



Fig. 1



Fig. 2



Fig. 3

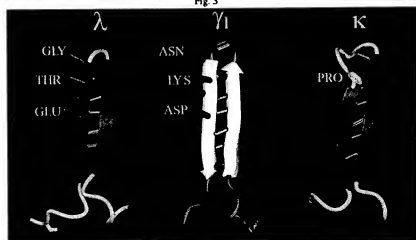


Fig. 4

Figs. 1–4.

chain (L-H) disulfide bond by the same distance as the comparable residues in the λ -type L chains of B7-15A2, New and Mcg.

10. Hydrogen bonding between symmetry-related κ -type V_L domains

As in strand 3–3 of C_L (see later section), strand 4–1 of the Pot V_L becomes ordered and relatively straight after a disruptive proline (residue 8). This ordered region consists of residues 9–13 (Ser–Ser–Leu–Ser–Ala), which are held in position on their inner surfaces by five hydrogen bonds with a parallel β -strand (5–5): Ser 9, Leu 11 and Ala 13 interact with residues Lys 103, Asp 105 and Lys 107. Along the outer surfaces of this strand, at least five carbonyl and amide backbone groups are appropriately posi-

tioned for intermolecular, antiparallel β -sheet formation with segment L9–13 in an adjacent Fv. In the center there are two hydrogen bonds (3.2 Å) between the backbone oxygen atoms of Ser L10 of Fv molecules 1 and 2 and the amide groups of Ser L12 of Fv molecules 2 and 1. However, the other two combinations of atoms are too far apart (3.6 Å) to complete the expected set of four hydrogen bonds. A stereo diagram illustrating these relationships is presented in Fig. 3.

11. Programming crystallization into the design of some Fabs

It may be possible to design human Fabs with features predisposing them to crystallize in accord with the packing principles discussed above. Five

Fig. 1. Top panel: Stereo diagram of three human B7-15A2 Fabs taken from a ribbon of molecules in the crystal lattice. The models were constructed and displayed with the program MOLMOL [10]. V domains are facing upward and the C domains downward in the outer two molecules. In the middle Fab the C domains are at the top. Polypeptide backbone is represented by tubular segments, yellow for the λ -type light (L) chain and steel blue for the γ 1-type heavy (H) chain. Strands of β -pleated sheets are shown as directional arrows superimposed on the polypeptide backbone. In the L chain, the 5-stranded layer in V_L and the 3-stranded layer in C_L are colored magenta; the four chain layers in V_L and C_L are in cyan. In the H chain, the 5- and 3-stranded layers are white and the 4-stranded layers are red. Hydrogen bonds stabilizing the packing of antiparallel molecules are represented by four purple rods between β -strands 3–3 of C_H1 and C_L (white to magenta on the right; magenta to white on the left). This intermolecular pattern of hydrogen bonds is repeated indefinitely along the ribbon in the crystal. Lower panel: Stereo diagram of three human (Mcg) λ -type Benze–Jones (L chain) dimers, excerpted as above from a ribbon of molecules in the crystal lattice. The monomers have the same amino acid sequence, but one behaves structurally as a heavy chain analog [11]. General orientations and color codes are the same as those in the upper panel. Intermolecular interactions include four hydrogen bonds on each side of the dimer, but in a slightly different distribution from that seen in B7-15A2 (see text and Table 1).

Fig. 2. Stereo diagram of a dimer of the Mcg \times Hud hybrid in the crystal lattice [12–14]. Mcg plays the structural role of the L chain and Hud is the H chain analog (both are λ chains). Note that the four intermolecular hydrogen bonds involve only the C_L domains of the Mcg component (purple rods crossing the crystallographic 2-fold axis between the two magenta 3–3 β -strands).

Fig. 3. Stereo diagram of a dimer of a human (Pot) Fv from an IgM- κ cryoglobulin. The κ -type V_L domain is represented by a yellow tubular model, with a cyan 4-chain layer and a magenta 5-chain layer. V_H is steel blue, with red (4-stranded) and white (5-chain) β -pleated sheets. The intermolecular hydrogen bonding interactions across a 2-fold axis involve the L chains only. Note that only the middle two sets of backbone atoms in the 4–1 β -strands are sufficiently close to fit the distance criteria (separation < 3.2 Å apart) for hydrogen bonds. The outer two sets of atoms, which are in the correct orientations for hydrogen bonds but 3.6 Å apart, are shown as small purple knots.

Fig. 4. Schematic representations of the last two strands of β -pleated sheets and their adjoining segments in the C domains of λ , γ 1 and κ chains. β -strand 3–2 is on the right and β -strand 3–3 is on the left of each drawing. Hydrogen bonds stabilizing the pleated sheets along their internal surfaces are depicted as ladders of purple rods between strands 3–2 and 3–3. Triads of amino acid residues involved in packing interactions of λ and γ 1 chains are listed next to their contributions to the external hydrogen bonds. In κ chains, a proline residue in a sequence position corresponding to threonine in λ chains and lysine in γ 1 chains interrupts the β -structure of strand 3–3. This disturbance leads to enlargement of the loop between strands 3–2 and 3–3 and a diminution or loss of the capacity to form β -pleated sheets with symmetry-related molecules. In λ chains, the connection between strands 3–2 and 3–3 is a narrow and regular β -turn. Two additional residues in the sequence of CH1 in γ 1 chains (Table 1) are accommodated by formation of a single turn of β_{10} helix at the crown of the loop. Thus the connection between strands 3–2 and 3–3 is kept relatively compact in γ 1 chains. In contrast to κ chains, the λ and γ 1 turns do not interfere with the propensity of the ensuing β -strand to pack with its neighbors in the crystal lattice.

human immunoglobulin fragments crystallizing primarily by antiparallel stacking of C domains (Mcg, Loc and Mcg \times Hud L chain dimers; New and B7-15A2 Fabs) all have λ -type L chains, alone or in combination with γ -type H chains of subclass I. It therefore seems prudent to emphasize this combination of isotypes in the initial attempts to program crystallization propensity.

A partial list of structural features making the λ and γ 1 isotypes so favorable for crystallization can be assembled from the previous discussions. (1) The 3–3 β -strands in the C domains are held in appropriate orientations by quite regular intra-domain hydrogen bonding. (2) Immediately following the turn between the 3–2 and 3–3 strands, the carbonyl oxygen atoms of the first residue (i) in 3–3 (Gly 203 in C_L and Asn 208 in C_H1) point outward for interactions with a docking molecule. (3) Backbone amide and carbonyl groups of residues $i+2$ and $i+4$ are also readily available for intermolecular hydrogen bonding. Among the known examples, they are the most frequently used entities for such purposes (e.g. Thr 205 and Glu 207 in C_L; Lys 210 and Asp 212 in C_H1). (4) Although there are small variations among the participants, four intermolecular hydrogen bonds between backbone atoms have been recorded in all five proteins of the λ and/or γ 1 isotypes. Interactions between side chains may add one-two hydrogen bonds to this count. (5) Side chains of the 3–3 β -strands do not interfere with the docking of the macromolecules. (6) The sets of i , $i+2$ and $i+4$ residues in the L and H chains are equidistant from the L-H interchain disulfide bond [9], which anchors the 3–3 β -strands and essentially terminates the Fab. This arrangement places the L and H β -strands in register for typical antiparallel hydrogen bonding. (7) Turns connecting the 3–2 and 3–3 β -strands are quite regular in the two isotypes and consequently do not significantly distort the 3–3 docking surfaces.

12. Exceptions to the rules: C_L domains of κ -type L chains

In two murine Fabs with κ chains (4-4-20 anti-

fluorescein and BV04-01 anti-ss-DNA antibodies), as well as the Human Pot Fv, we found that the L chains dominated the crystal packing but not by the mechanisms emphasized here [4–8]. Fragments with κ -type L chains are still missing from our preliminary list of proteins predisposed to crystallize by C domain stacking. A partial explanation may lie in the structure of the turn between β -strands 3–2 and 3–3 of a κ -type C_L domain in a humanized (AF2) Fab (Z.-C. Fan and A.B. Edmundson, unpublished results). This turn is distinctly different from those of λ and γ 1 chains of B7-15A2, as illustrated in Fig. 4. The more regular turns of λ and γ 1 chains will be discussed first.

13. Structures of turns between 3–2 and 3–3 in the B7-15A2 Fab

Amino acid sequences of both the human κ and γ 1 chains have two more residues than λ chains between the intrachain and interchain disulfide bonds [9], as shown in Table 1. A gap is introduced into the λ chain sequence to indicate the locations of these extra residues, which are probably the ones that expand the β -turn in the λ chain to a distorted helix in the γ 1 chain.

On the amino side of the turn (strand 3–2), all three polypeptide chains are very similar (see Fig. 4). In the λ chains of both B7-15A2 and Mcg, the connecting segment between 3–2 and 3–3 is a compact type II β -turn of four residues (His–Glu–Gly–Ser, nos. 201–204). This turn is kept narrow and well ordered by two hydrogen bonds between His 201 and Ser 204 (i , $i+3$) at its base.

Table 1
COOH-terminal amino acid sequences of human λ , γ 1 and κ chains

CQVTHE	GSTVEKTVAPTECS	λ Chain
CNVNHKPSNTKVDKRVEPKSC		γ 1 Chain
CEVTHQGLSSPVTKSFNRGE C		κ Chain

The abbreviations are presented in the one-letter code. Symbols in boldface type represent residues involved in packing interactions in λ and γ 1 chains, and their counterparts in κ chains.

Moreover, this arrangement insures a stable platform for the carbonyl oxygen atom of Gly 203 to participate in external hydrogen bonding with another L or H chain.

In the B7-15A2 H chain, the turn is more irregular and slightly wider than its counterpart in the λ chain. For instance, the loop is stabilized at its base by a single hydrogen bond between His 210 and Thr 215 ($i, i+5$), which demarcate where β -strand 3–2 ends and 3–3 begins. The crown of the loop (Lys–Pro–Ser–Asn, nos. 211–214) flares outward between these two positions to form a single turn of a distorted 3_{10} helical structure, with hydrogen bonding between Lys 211 and Asn 214 (see crest of the loop in Fig. 4). At the end of the loop, the backbone carbonyl group of Asn 214 is in an orientation suitable for intermolecular hydrogen bonding. Structurally, Asn 214 plays a roll like that of Gly 203 of the λ chain.

14. Structure of the κ chain loop between β -strands 3–2 and 3–3

Two more residues are taken from the generic strand 3–3 and added to produce the κ chain loop, which then has a combined sequence of His–Gln–Gly–Leu–Ser–Ser–Pro–Val (residues 198–205). Thus the λ , $\gamma 1$ and κ loops progress in length from four to six to eight residues. The conformation of β -strand 3–2 of the κ chain is similar to that of the $\gamma 1$ chain through the His residue. After the turn, the κ polypeptide chain comes back into register with its $\gamma 1$ counterpart at Val 205. From Gln 199 to Pro 204 the segments are very different, as is evident from Fig. 4.

Note that the loop is considerably broader and more irregular than those of the λ and $\gamma 1$ chains. Moreover, residues predicted to be potential contact sites for other L or H chains in the crystal lattice (e.g. Gly 200, Ser 202 and Pro 204) are constituents of the looped out structure, rather than the relatively straight part of β -strand 3–3.

It is now clear that the ordered (β -strand) portion of 3–3 does not begin in a κ chain until Val 205, which corresponds structurally to the center of the crystal packing surfaces of the λ and $\gamma 1$ chains. The residue most responsible for this

anomaly is Pro 204, which not only alters the direction of the chain but also disrupts the hydrogen bonding propensity (only the carbonyl oxygen of proline can participate).

By sequence criteria, Ser 202, Pro 204 and Thr 206 in a κ chain are equivalent to λ constituents Gly 203, Thr 205 and Glu 207, the three most commonly used residues for stacking interactions in crystals. Because of the complementarity requirements, we suspect that the surface on the carboxyl side of Pro 204 would not be sufficiently large or topographically adequate to bind a C_L or C_{H1} domain in a symmetry-related Fab.

15. Usage of amino acid triads in C_L – C_L or C_L – C_{H1} packing

β -strand 3–3 begins with the carbonyl group of glycine in λ chains (at the end of the gap in the sequences given in Table 1) and with asparagine in a comparable position in $\gamma 1$ chains. These are the first of three residues (boldface type in Table 1) involved in cross-molecule hydrogen bonding in human λ and $\gamma 1$ chains. Corresponding residues in the κ chain are also in boldface. As emphasized earlier, the first two are constituents of the turn between strands 3–2 and 3–3.

Note that the residues in the packing triads of λ and $\gamma 1$ chains are in register relative to the inter-chain disulfide bond and are therefore easy to stack in an antiparallel fashion. The first residue in the triad only supplies its carbonyl group to the intermolecular interactions but the backbone amide and carbonyl groups of the other two members are fully accessible.

When inserted into the middle of what would otherwise be a packing triad in C_L of a κ chain, proline disrupts the type of ordered local structure seen in the λ and $\gamma 1$ chains. The relatively straight stretch of the 3–3 β -strand is thereby converted into part of a wide turn less suitable for antiparallel docking of another C domain. A proline residue also interrupts the order of β -strand 4–1 in the Pot V_L domain. Because of these disruptive effects, as well as changes resulting from the parallel pattern of internal hydrogen bonding, the packing triad for the Pot Fv is Ser 9, Ser 10 and

Ser 12, rather than the more regular $i, i+2, i+4$ arrangement of residues in C_L and C_H1 .

16. Conclusion

For human antibodies with λ and $\gamma 1$ chains, it seems straightforward to program a crystallization tendency into the design of synthetic Fabs. We can follow the leads provided by X-ray analyses of Fabs from naturally occurring antibodies. In these molecules, the 3–3 β -strands and the turns between strands 3–2 and 3–3 in both C_L and C_H1 are highly conserved in length and sequence.

From a three-dimensional viewpoint, crystal packing of C_L and C_H1 seems to be more dependent on the presence of a highly ordered 3–3 β -strand than on the actual sequence of side chains. For example, the packing triads consist of Gly, Thr and Glu residues in the λ chains and Asn, Lys and Asp residues in the $\gamma 1$ chains. As a generalization, the side chains do not interfere with the packing interactions and may even enhance them (e.g. Mcg L chain). However, a single substitution (proline) in the triad region of a κ chain can disrupt the structural order of the 3–3 β -strand and render it unsuitable for stacking with another C_L or C_H1 . It would be interesting to see if the structure of the 3–3 strand could be significantly altered by replacement of the proline.

In Fabs with $\gamma 1$ chains, the last six residues (including the interchain Cys) of the H chain are encoded by the 'hinge' region gene. Different subclasses of human and murine IgG antibodies (as well as IgM and IgA immunoglobulins) have distinct hinge regions with respect to disulfide patterns, sequences and length. During the early tests of the ideas presented here, it is therefore desirable to program $\gamma 1$ -type hinge regions into synthetic Fabs. This decision will help insure that the packing triads of C_L and C_H1 are kept in register relative to the interchain disulfide bond.

Fabs with κ -type L chains are not expected to crystallize as ribbons but there is no a priori reason why their H chain ($\gamma 1$) components

should not pack as dimers. For Fvs with κ chains (e.g. Pot), dimerization through the formation of cross molecule β -sheet structures can produce high quality crystals with favorable protein to solvent ratios.

In the Pot Fv, side chains are involved in the cross-molecule interactions and the sequence in this part of framework region 1 (FR1) is therefore of considerable importance. Of the three subgroups of human κ chains [9], only subgroup II has an unfavorable sequence (a proline in position 12) for the Pot type of packing. Five side chains of strand 4–1 in Pot are in contact with a symmetry related L chain and it seems remarkable that they all dovetail properly.

This information should be of substantial value in the synthesis and crystallization of synthetic Fvs and single chain antibodies with κ -type L chains. In our limited data base, the κ chains are also prominent in controlling the packing of larger antibody fragments, although not as yet in predictable ways. Both λ and $\gamma 1$ chains have naturally occurring features that need little embellishment to make them self assemble into highly ordered patterns. These patterns give great delight to contemporary crystallographers and molecular biologists who are trying to extend the new field of structure based design to the production of synthetic antibodies.

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COMMUNICATION

SCOP: A Structural Classification of Proteins Database for the Investigation of Sequences and Structures

Alexey G. Murzin, Steven E. Brenner, Tim Hubbard and Cyrus Chothia*

MRC Laboratory of Molecular
Biology and Cambridge
Centre for Protein
Engineering, Hills Road
Cambridge CB2 2QH
England

To facilitate understanding of, and access to, the information available for protein structures, we have constructed the Structural Classification of Proteins (scop) database. This database provides a detailed and comprehensive description of the structural and evolutionary relationships of the proteins of known structure. It also provides for each entry links to co-ordinates, images of the structure, interactive viewers, sequence data and literature references. Two search facilities are available. The homology search permits users to enter a sequence and obtain a list of any structures to which it has significant levels of sequence similarity. The key word search finds, for a word entered by the user, matches from both the text of the scop database and the headers of Brookhaven Protein Databank structure files. The database is freely accessible on World Wide Web (WWW) with an entry point to URL <http://scop.mrc-lmb.cam.ac.uk/scop/>

scop: an old English poet or minstrel (Oxford English Dictionary);

oxon: pile, accumulation (Russian Dictionary).

Keywords: protein families; superfamilies; folds; evolutionary relationships

*Corresponding author

Nearly all proteins have structural similarities with other proteins and, in many cases, share a common evolutionary origin. The knowledge of these relationships makes important contributions to molecular biology and to other related areas of science. It is central to our understanding of the structure and evolution of proteins. It will play an important role in the interpretation of the sequences produced by the genome projects and, therefore, in understanding the evolution of development.

The recent exponential growth in the number of proteins whose structures have been determined by X-ray crystallography and NMR spectroscopy means that there is now a large and rapidly growing corpus of information available. At present (January, 1995) the Brookhaven Protein Databank (PDB, (Abola *et al.*, 1987)) contains 3091 entries and the number is increasing by about 100 a month. To facilitate the understanding of, and access to, this information, we have constructed the Structural Classification of Proteins (scop) database. This database provides a detailed and comprehensive description of the structural and evolutionary relationships of proteins whose three-dimensional structures have been determined. It includes all

proteins in the current version of the PDB and almost all proteins for which structures have been published but whose co-ordinates are not available from the PDB.

The classification of protein structures in the database is based on evolutionary relationships and on the principles that govern their three-dimensional structure. Early work on protein structures showed that there are striking regularities in the ways in which secondary structures are assembled (Levitt & Chothia, 1976; Chothia *et al.*, 1977) and in the topologies of the polypeptide chains (Richardson, 1976, 1977; Sternberg & Thornton, 1976). These regularities arise from the intrinsic physical and chemical properties of proteins (Chothia, 1984; Finkelstein & Pitts, 1987) and provide the basis for the classification of protein folds (Levitt & Chothia, 1976; Richardson, 1981). This early work has been taken further in more recent papers; see, for example, Holm & Sander (1993), Orengo *et al.* (1993), Overington *et al.* (1993) and Yee & Dill (1993). An extensive bibliography of papers on the classification and the determinants of protein folds is given in scop.

The method used to construct the protein classification in scop is essentially the visual inspection and comparison of structures through various automatic tools are used to make the task manageable and help provide generality. Given the

Abbreviations used: PDB, Protein Databank; scop, Structural Classification of Proteins.

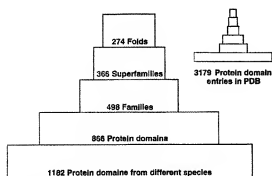


Figure 1. In scop, the unit of classification is usually the protein domain. Small proteins, and most of those of medium size, have a single domain and are, therefore, treated as a whole. The domains in large proteins are usually classified individually. The protein entries in the December 1994 of the Brookhaven Protein Databank (PDB) contain 3179 domains. Many of these become forms of the same protein whose differences are not significant in terms of the classification used here; for example they have different bound ligands or engineered mutations. To distinguish between these and structures of the same protein from different organisms, proteins listed within a family are subclassified by species. Classification of the 3179 domains show that they come from 498 families that is clustered into 366 superfamilies and 279 different folds. In addition to these, scop contains entries for 195 proteins that do not have atomic co-ordinates available from the PDB at present but for which description of their structures have been published.

current limitations of purely automatic procedures, we believe this approach produces the most accurate and useful results. The unit of classification is usually the protein domain. Small proteins, and most of those of medium size, have a single domain and are, therefore, treated as a whole. The domains in large proteins are usually classified individually.

The classification is on hierarchical levels that embody the evolutionary and structural relationships.

FAMILY. Proteins are clustered together into families on the basis of one of two criteria that imply their having a common evolutionary origin: first, all proteins that have residue identities of 30% and greater; second, proteins with lower sequence

identities but whose functions and structures are very similar; for example, globins with sequence identities of 15%.

SUPERFAMILY. Families, whose proteins have low sequence identities but whose structures and, in many cases, functional features suggest that a common evolutionary origin is probable, are placed together in superfamilies; for example, actin, the ATPase domain of the heat-shock protein and hexokinase (Flaherty *et al.*, 1991).

COMMON FOLD. Superfamilies and families are defined as having a common fold if their proteins have same major secondary structures in same arrangement with the same topological connections. In scop we give for each fold short descriptions of its main structural features. Different proteins with the same fold usually have peripheral elements of secondary structure and turn regions that differ in size and conformation and, in the more divergent cases, these differing regions may form half or more of each structure. For proteins placed together in the same fold category, the structural similarities probably arise from the physics and chemistry of proteins favouring certain packing arrangements and chain topologies (see above). There may, however, be cases where a common evolutionary origin is obscured by the extent of the divergence in sequence, structure and function. In these cases, it is possible that the discovery of new structures, with folds between those of the previously known structures, will make clear their common evolutionary relationship.

CLASS. For convenience of users, the different folds have been grouped into classes. Most of the folds are assigned to one of the five structural classes on the basis of the secondary structures of which they composed: (1) all alpha (for proteins whose structure is essentially formed by α -helices), (2) all beta (for those whose structure is essentially formed by β -sheets), (3) alpha and beta (for proteins with α -helices and β -strands that are largely interspersed), (4) alpha plus beta (for those in which α -helices and β -strands are largely segregated) and (5) multi-domain (for those with domains of different fold and for which no homologues are known at present). Note that we do not use Greek characters in scop because they are not accessible to all world wide web viewers. More unusual proteins, peptides and the PDB entries for designed proteins,

Table 1

Facilities and databases to which SCOP has links

Link	Source	URL	Reference
Co-ordinates	PDB	http://www.pdb.bnl.gov/	(Abola <i>et al.</i> , 1987)
Static Images	SP3D	http://expasy.chugce.ch/gopher://pdb.pdb.bnl.gov/	(Appel <i>et al.</i> , 1994)
On-the-fly images	NIH molecular modelling group	http://www.nih.gov/www94/molvis	(FitzGerald, 1994)
Sequences and MEDLINE entries	NCBI Entrez	http://www.ncbi.nlm.nih.gov/	(Benson <i>et al.</i> , 1993)

The scop database contains links to a number of other facilities and databases in the world. Several interactive viewers can be linked with scop using PDB co-ordinates. The location and nature of the links will vary as databases evolve and relocate.

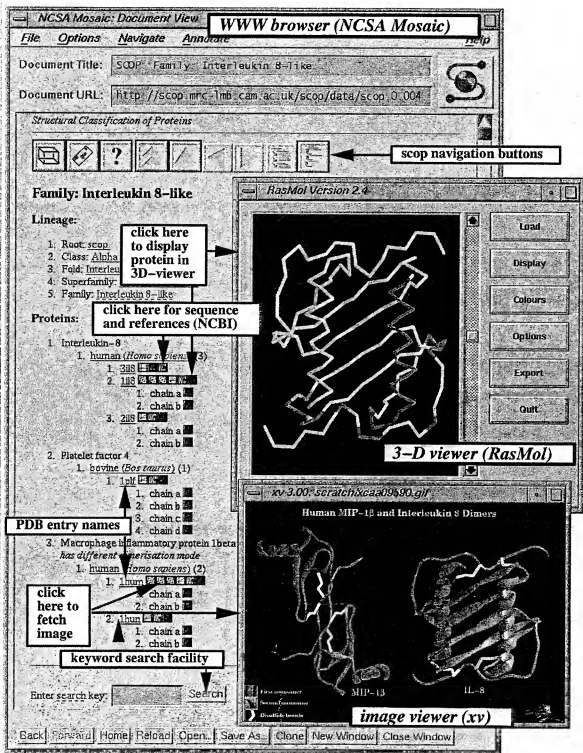


Figure 2. A typical scop session is shown on a unix workstation. A scop page, of the Interleukin 8-like family, is displayed by the WWW browser program (NCSA Mosaic) (Schatz & Hardin, 1994). Navigating through the tree structure is accomplished by selecting any underlined entry by clicking on buttons (at the top of each page) and by keyword searching (at the bottom of each page). The static image comparing two proteins in this family was downloaded by clicking on the icon indicated and is displayed by image-viewer program xv. By clicking on one of the green icons, commands were sent to a molecular viewer program (RasMol) written by Roger Sayle (Sayle, 1994), instructing it to automatically display the relevant PDB file and colour the domain in question by secondary structure. Since sending large PDB files over the network can be slow, this feature of scop can be configured to use local copies of PDB files if they are available. Equivalent WWW browsers, image-display programs and molecular viewers are also available free for Windows-PC and Macintosh platforms.

theoretical models, nucleic acids and carbohydrates, have been assigned to other classes.

The number of entries, families, superfamilies and common folds in the current version of scop are shown in Figure 1. The exact position of boundaries between family superfamily and fold are, to some degree, subjective. However, because all proteins that could conceivably belong to a family or superfamily are clustered together in the encompassing fold category, some users may wish to concentrate on this part of the database.

In addition to the information on structural and evolutionary relationships, each entry (for which co-ordinates are available) has links to images of the structure, interactive molecular viewers, the atomic co-ordinates, sequence data and homologues and MEDLINE abstracts (see Table 1).

Two search facilities are available in scop. The homology search permits users to enter a sequence and obtain a list of any structures to which it has significant levels of sequence similarity. The key word search finds, for a word entered by the user, matches from both the text of the scop database and the headers of Brookhaven Protein Databank structure files.

To provide easy and broad access, we have made the scop database available as a set of tightly coupled hypertext pages on the world wide web (WWW). This allows it to be accessed by any machine on the internet (including Macintoshes, PCs and workstations) using free WWW reader programs, such as Mosaic (Schatz & Hardin, 1994). Once such a program has been started, it is necessary only to "open" URL.

<http://scop.mrc-lmb.cam.ac.uk/scop/>

to obtain the "home" page level of the database.

In Figure 2 we show a typical page from the database. Each page has buttons to go back to the top-level home page, to send electronic mail to the authors, and to retrieve a detailed help page. Navigating through the tree structure is simple; selecting any entry retrieves the appropriate page. In addition, buttons make it possible to move within the hierarchy in other manners, such as "upwards" to obtain broader levels of classification.

The scop database was originally created as a tool for understanding protein evolution through sequence-structure relationships and determining if new sequences and new structures are related to previously known protein structures. On a more general level, the highest levels of classification provide an overview of the diversity of protein structures now known and would be appropriate both for researchers and students. The specific lower levels should be helpful for comparing individual structures with their evolutionary and structurally related counterparts. In addition, we have also found that the search capabilities with easy access to data and images make scop a powerful general-purpose interface to the PDB.

As new structures are released by PDB and published, they will be entered in scop and revised

versions of the database will be made available on WWW. Moreover, as our formal understanding of relationships between structure, sequence function and evolution grows, it will be embodied in additional facilities in the database.

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Complex between *Peptostreptococcus magnus* Protein L and a Human Antibody Reveals Structural Convergence in the Interaction Modes of Fab Binding Proteins

Marc Graille,¹ Enrico A. Stura,^{1,2} Nicholas G. Housden,² Jennifer A. Beckingham,² Stephen P. Bottomley,² Dennis Beale,³ Michael J. Taussig,³ Brian J. Sutton,⁴ Michael G. Gore,² and Jean-Baptiste Charbonnier^{1,2}

¹Département d'Ingénierie et d'Etudes des Protéines
Commissariat à l'Energie Atomique
Centre d'Etudes Saclay
Gif-sur-Yvette F-91191
France

²Department of Biochemistry
Institute of Biomolecular Sciences
University of Southampton
Bassett Crescent East
Southampton SO16 7PX
United Kingdom

³Laboratory of Molecular Recognition
The Babraham Institute
Babraham, Cambridge CB2 4AT
United Kingdom

⁴The Randall Centre
King's College London
Guy's Campus
London SE1 1UL
United Kingdom

Summary

Background: *Peptostreptococcus magnus* protein L (PpL) is a multidomain, bacterial surface protein whose presence correlates with virulence. It consists of up to five homologous immunoglobulin binding domains that interact with the variable (V_L) regions of kappa light chains found on two thirds of mammalian antibodies.

Results: We refined the crystal structure of the complex between a human antibody Fab fragment (2A2) and a single PpL domain (61 residues) to 2.7 Å. The asymmetric unit contains two Fab molecules sandwiching a single PpL domain, which contacts similar V_L framework regions of two light chains via independent interfaces. The residues contacted on V_L are remote from the hypervariable loops. One PpL-V_L interface agrees with previous biochemical data, while the second is novel. Site-directed mutagenesis and analytical-centrifugation studies suggest that the two PpL binding sites have markedly different affinities for V_L. The PpL residues in both interactions are well conserved among different *Peptostreptococcus magnus* strains. The Fab contact positions identified in the complex explain the high specificity of PpL for antibodies with kappa rather than lambda chains.

Conclusions: The PpL-Fab complex shows the first interaction of a bacterial virulence factor with a Fab light chain outside the conventional combining site. Struc-

tural comparison with two other bacterial proteins interacting with the Fab heavy chain shows that PpL, structurally homologous to streptococcal SpG domains, shares with the latter a similar binding mode. These two bacterial surface proteins interact with their respective immunoglobulin regions through a similar β zipper interaction.

Introduction

Peptostreptococcus magnus protein L (PpL: whole protein L, PpL domain: individual domain of protein L) is a cell wall-anchored protein able to interact with a large repertoire of mammalian immunoglobulins (Ig) [1]. *Staphylococcus aureus* protein A (SpA) [2] and streptococcal protein G (SpG) [3] also share this property, although they recognize different Ig regions. PpL is present at the surface of about 10% of *Peptostreptococcus magnus* strains [4]. It is a 76–106 kDa protein containing four or five highly homologous, consecutive extracellular Ig binding domains (depending on the bacterial strain from which it is isolated [5, 6]). The structure of PpL domain B (78 amino acids) has been determined by NMR spectroscopy [7]. The fold of this domain is similar to that of the SpG Ig binding domains. It consists of a β sheet composed of two pairs of anti-parallel β strands and an α helix that lies on top of the sheet.

PpL interacts with Ig light chains [1], notably with the kappa light-chain variable region (V_L) from humans and other mammals [8, 9]. When present on the bacterial surface, PpL has been described as a virulence factor of bacterial vaginosis in different clinical specimens [4]. It has also been shown that PpL induces histamine release by basophils and mast cells, presumably by cross-linking IgE bound to Fc ϵ receptors [10, 11].

PpL and SpA single domains are targets for protein engineering due to their ability to bind the variable regions (Fv) of a large population of antibodies. This unique property makes them valuable tools in biotechnology for the purification and recognition of recombinant single-chain (sc) Fv. Such engineered antibodies are increasingly used in the optimization of specificity and affinity by phage display and other *in vitro* evolutionary mutagenesis techniques [12]. The structural characterization of PpL and SpA binding properties is useful for defining the spectrum of Fvs, which bind to these domains, based on their sequences.

We report in this study the crystal structure of the human antibody Fab 2A2 complexed through its V_L region to a PpL domain. The same Fab was the subject of a previous crystallographic study describing its complex with a *Staphylococcus aureus* protein A (SpA) domain, which binds to the antibody V_L region [13]. Here, we compare the location of these two Ig binding domains on either side of the Fv region of Fab 2A2 in relation to the antigen-combining site. Unexpectedly, we find that

¹ Correspondence: estura@cea.fr (E.A.S.),
jb.charbonnier@cea.fr (J.B.C.)

Key words: immunoglobulin binding proteins (IBP); antibody complex; cell surface protein; X-ray structure

there are two PpL-Fab interfaces, such that one PpL domain has two separate regions that can interact with kappa light chains and is capable of binding two Fabs simultaneously. The two interfaces involve similar sites on the V_L domains. One of the PpL-Fab interfaces conforms to previous biochemical data, while the second is novel. Site-directed mutagenesis and analytical-centrifugation studies suggest that the two PpL combining regions have markedly different affinities for V_L . The high specificity of PpL for kappa (the largest mammalian V_L gene family), rather than lambda chains, is discussed in light of the structure of the complex. We analyze the sequence diversity of the PpL domains at positions involved in the interaction with V_L regions and discuss avidity effects reported between whole PpL and Ig. Furthermore, by comparing the interactions of PpL and SpG with V_L and G_1 , respectively, we note that at a structural level, the Fab binding modes of these two bacterial proteins show a degree of convergence.

Results and Discussion

The First V_L -PpL Interface

The asymmetric unit has one PpL domain and two Fab molecules. The PpL domain is in close contact with the V_L region from both Fabs (Figure 1a). This stoichiometry was unexpected because no previous studies raised any evidence for one PpL domain being able to complex to two V_L regions simultaneously [6, 14].

The first interface buries a total solvent-accessible area of 1300 Å² with approximately equal contributions from both molecules, as determined with a 1.4 Å radius probe, and is remote from the Ig heavy chain. There is no conformational change in the backbone of either partner upon binding. The PpL domain C* used in this study has 94% sequence identity with the C4 domain [15]. Its affinity for kappa chains is comparable (130 nM) to the one measured for PpL domain B1 [14, 15]. Domain C* maintains the same three-dimensional structure as PpL domain B1 determined by NMR [7], and the two structures superpose with an rmsd of 1.31 Å over 59 residues. The complexed PpL domain C* superposes with an rmsd of 0.39 Å over 61 residues of the crystallographic structure of free PpL domain C* (B.J.S., unpublished data). Similarly, the variable region of the complexed Fabs superimposes with an rmsd of 0.6 Å over 215 residues with the free structure determined previously [15].

The interaction with PpL involves 13 residues from the Fab (Figures 1b and 1c). Ten are located in framework region 1 (FR1). The others are Lys-L107 from the segment connecting the V_L to the C₁ region, Glu-L143 from the C₁ region, and Arg-L24 from the CDR-L1 region [16] on strand B. However, this residue does not belong to the V_L hypervariable loops according to the structural definition of Chothia [17] or to the positions frequently identified by the contact with antigens [18]. Compared with SpA, which binds to the variable region of the same human antibody (Fab 2A2) [13], PpL is farther away from the center of the antigen binding site (23 Å compared to 16 Å). Hence, like SpA, PpL binding should not affect accessibility to the antigen-combining site, as also suggested by competition assays [19].

Of the amino acids in the PpL domain, 12, located mainly on strand $\beta 2$ and the α helix, are involved in this interaction (Figure 2). This interface is characterized by six hydrogen bonds (Figure 1b, Table 1). Three are between main-chain atoms located on PpL strand $\beta 2$ and on strand A from the first Fab; they thus join the β sheets of the Fab and the protein L into a unique sheet through a β zipper interaction.

Heteronuclear NMR spectroscopy has been used for mapping backbone positions of the PpL domain B1 [20] involved in the interaction with the V_L region. Most positions identified for domain B1 are also implicated in the first interface of domain C* described in the present study. These backbone positions are on strand $\beta 2$ and the α helix of the PpL domains. A minor discrepancy with the NMR results concerns the loop between the α helix and strand $\beta 3$, which does not make any contact with the Fab V_L region in the crystal structure of the complex. This loop is poorly defined in the NMR structure, so the discrepancy can be attributed to a change in mobility upon complexation.

The first interface was subjected to site-directed mutagenesis of PpL domain C*. A 23-fold drop in affinity results from a Y53P substitution [21] as measured by competitive ELISA. This decrease following the loss of the tyrosine hydroxyl is as expected on the basis of the energy of a neutral hydrogen bond [22], consistent with the loss of the interaction between the Tyr-53 hydroxyl group and the Thr-L20 carbonyl group of the V_L region (Table 1). Thus, the first interface explains the existing data well.

The Second V_L -PpL Interface

In the crystal, the single PpL domain is sandwiched between the V_L regions of two 2A2 Fabs present in the asymmetric unit (Figure 1a). The second interaction buries a total solvent-accessible area of 1400 Å², comparable to other protein-protein complexes [23]. The PpL domain C* has a different orientation in the two interactions relative to the Fab β sheet. This second interaction involves 15 V_L residues, located mainly on the β strands A and B (as in the first interface) with some participation of strands D and E (Figure 1c). Out of the 15 residues from the V_L region involved in this second interaction, 10 are common to the first one. On the contrary, none of the PpL residues that contribute significantly to this interface are involved in the first one. The 14 amino acids from the PpL domain involved in the second interaction come mainly from strand $\beta 3$ and from the α helix (Figure 2). Sixtyhydrogen bonds and two salt bridges mediate this interaction (Table 2), as compared to only six hydrogen bonds for the first interface (Table 1). Although unexpected from biochemical studies, this second PpL-Fab interface buries an area too large to be a crystal contact. Given that this interface buries a surface larger than 1400 Å², the probability that it is just a crystal contact can be evaluated as only 2% [24]. Thus, we believe that this interface has to be given serious attention.

What is the strength of the second interaction? At present, no definite answer can be given, but we have used mutagenesis to probe the relative contributions of these two interfaces. First, we constructed the Y54W

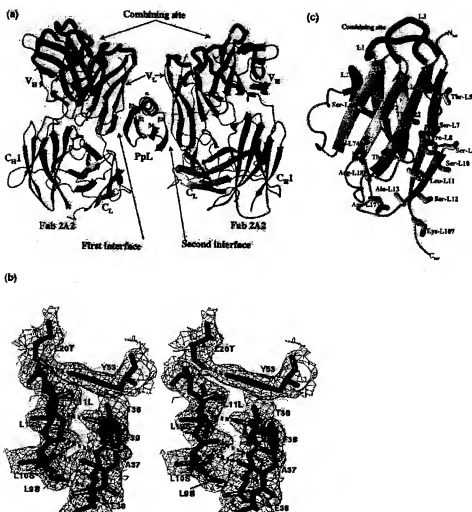


Figure 1. PpL domain C' Complexed with Human IgM Fab 2A2

(a) Ribbon representation of the 2 Fab:1 PpL domain complex present in the asymmetric unit. The PpL domain C' (red) is sandwiched between two Fab (blue and green). Light colors represent the light chains, and dark colors represent the heavy chains. Magenta highlights the CDR loops, as defined by Chothia [17], and positions the Fab-PpL interfaces relative to the combining site. Pseudo 2-fold axis symmetry relates the two Fabs, but both interfaces are not symmetrical.

(b) Stereo view of the 2F₅ - F₂ electron density map contoured at 1σ at the first V_H-PpL interface. The letter L precedes amino acids from the Fab light chain. Green dotted lines depict the three hydrogen bonds between Fab and PpL main-chain atoms. The three other hydrogen bonds are shown in magenta and involve at least one side-chain atom.

(c) Ribbon representation of the V_L region of Fab that interacts with the PpL domain. Ten residues common to the first and second interfaces are in yellow. Positions in pink are only involved in the first interface, and those in light green are implicated only in the second interface. All figures were generated with MOLMOL [42] or TURBO-FRODO [43].

mutant of a PpL domain C' to have an efficient fluorescent probe so as to measure K_d by the stopped-flow method [21]. Second, we constructed two double mutants, Y53F-Y64W [21] and D55A-Y64W (this study), with the purpose of weakening, respectively, the first and

second interfaces. In the Y53F-Y64W mutant, the replacement of Tyr-53 by Phe disrupts the hydrogen bond between the tyrosine hydroxyl and the L20 carbonyl in the first interface. Similarly, we chose the D55A-Y64W mutant in order to disrupt the salt bridge between Asp-

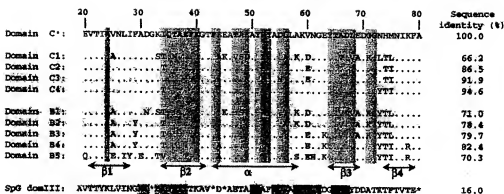


Figure 2. Alignment of the Amino Acid Sequences of the Different Ig Binding Domains from PpL.

The PpL residues represented are only those visible in the crystal structure. Domain C' is the one used in this study. Domains C, to C, are from PpL₂₀₀ strain [5], and domains B, to B, are from PpL₂₀₀ strain [6]. PpL residues involved in the first interface with the Fab 2A2 are highlighted in pink, and those involved in the second binding site are in green. With the exception of Arg-52 (yellow), there is no overlap between the residues involved in each interaction. Core residues are in bold. Those involved in the second interface are mainly located on the α helix and strand $\beta 3$ and are well conserved among the different PpL domains. Of note is that one mutation (Gly-71 to Lys) may reinforce or weaken the second interface. The amino acid sequence of SpG domain III is also shown with the residues involved in the interaction with the Fab shown in pink, those interacting with the Fc γ in cyan, and those implicated in both interactions in yellow.

55 and Arg-L24 as well as the hydrogen bond with the Ser-L7 side chain in the second interface. These three single PpL domain constructs are correctly folded, as indicated by far-UV CD spectroscopy (data not shown). While the loss of a single hydrogen bond at the first interface has a significant effect on the dissociation constant of the V_L-PpL domain C' complex, the disruption of the salt bridge and the hydrogen bond mediated by Asp-55 of PpL at the second interface does not alter the K_d of the complex (Table 3). This suggests that the binding of single PpL domain C' is dominated by the first interface and that the dissociation constant of the second is at least one order of magnitude larger than the first one.

Evidence supporting the existence on a single PpL domain of two Fab-combining regions, differing in affinities by about two orders of magnitude, has been obtained recently by analytical-centrifugation studies (B.J.S. and R. Beavil, unpublished data).

This observation is reminiscent of the 1:2 stoichiometry described for the interaction between human growth hormone and its receptor [25], where two different faces of the hormone contact similar surfaces on two recep-

tors, with the result being receptor activation [26]. By analogy, in our study a single PpL domain contacts similar surfaces on two V_L regions and could bridge two Igs anchored at the membrane of B cells. The biological relevance of this second interface and the potential mitogenic activity of a single PpL domain on B cells are under investigation.

V_L Specificity of the PpL Binding Interaction

PpL has been reported to bind efficiently to V_L regions of about two thirds of the human Ig repertoire and to thus encompass mainly the $\kappa 1$, $\kappa 3$, and $\kappa 4$ subgroups [8] but neither $\kappa 2$ nor any λ subgroups. The PpL kappa specificity observed for human antibodies extends to various other mammalian species [9]. To analyze the structural determinants of this specificity, we have superimposed on the 2A2 V_L region the equivalent structures found on different kappa and lambda subgroups. We observed that PpL binding ability is mostly concen-

Table 1. Hydrogen Bonds Involved in the First V_L-PpL Interface

V _L Region	PpL Domain C'	Distance (Å)
Ser L-9 N	O Glu 38	2.8
Ser L-9 Oy	N 40 Lys	3.1
Ser L-10 Oy	O ₁ Glu 38	2.9
Ser L-10 N	N Glu 38	3.1
Ser L-12 N	O Thr 38	2.7
Thr L-20 O	O ₁ Tyr 53	2.8

Hydrogen bond distances were calculated with the program CONTACTS [44] with a maximum distance cutoff of 3.4 Å and defined according to the criteria of McDonald and Thornton [45].

Table 2. Hydrogen Bonds and Salt Bridges Involved in the Second V_L-PpL Interface

V _L Region	PpL Domain C'	Distance (Å)
Ser L-7 Oy	O81 Asp 55	2.9
Ser L-10 Oy	O ₁ Thr 55	3.2
Ser L-12 N	O Ala 55	2.9
Ser L-12 O	N Leu 55	2.8
Arg L-18 N	O Gly 71	3.1
Arg L-24 N ₁	O81 Asp 55	2.7
Lys L-107 N ₂	O81 Asp 57	2.8
Lys L-107 N ₂	O Leu 55	2.9

Hydrogen bond and salt bridge distances were calculated with the program CONTACTS [44] with a maximum distance cutoff of 3.4 Å and defined according to the criteria of McDonald and Thornton [45].

Table 3. Dissociation Constants between PpL Single Domain C' and Human κ Chain as Determined by Stopped-Flow Fluorescence Measurements

PpL Construct	K _d (μ M)	Comment
Wild type	0.13 ¹	
Y84 W	0.17 ²	Fluorescent probe
Y83F-Y84W	2.8 ²	Hydrogen bond on interface 1 disrupted
D85A-Y84W	0.145	Hydrogen bond and salt bridge on interface 2 disrupted

¹From Beckingham et al. [14].

²From Beckingham et al. [21].

trated on the L5 to L12 segment of the V_L region and that the interaction is dependent on the main-chain conformation of this segment, on which conformation the β zipper interaction also depends. Particularly important are residues L8–L12, which bury more than 80% of their total accessible area. The V_L residues of this segment and at other positions of the first interface are well conserved among the recognized V_L regions (Table 4). The main chain superimposes well onto V_L structures of the κ 1, κ 3, and κ 4 subgroups (rmsd of 0.3–0.5 Å over the eight residues from L5 to L12). Most κ 2 subgroup sequences have a proline residue at position L12 that introduces major steric hindrance by pointing the Pro ring toward the PpL domain. For the λ subgroup, two observations account well for the absent or weak binding activity found for these chains. Firstly, the L5–L12 segment is one residue shorter, resulting in backbone conformational differences, which may alter the β zipper interaction (rmsd of 1–1.2 Å over the seven residues from L5 to L12). Secondly, due to the reduced size of this segment, a cavity is created around positions L7 and L8. The Fab recognition mode developed by the PpL domain is highly dependent on the backbone conformation, different from that of SpA, which relies more on a side chain-specific recognition mode [13].

Avidity of Whole PpL for Immunoglobulins

PpL contains four or five highly homologous, consecutive extracellular Ig binding domains in tandem, depending on the strain. At what level are the positions involved in the first interface conserved between the different domains? Sequence alignment of the four C

domains and five B domains with the C' domain (Figure 2) shows that domain C1, which differs at 20 positions out of 61, is the most distant from domain C'. The differences on C1 are distributed all over the domain, with higher incidence on the α helix and strands β 2, β 3, and β 4. The other C domains have a higher degree of sequence identity with C'. The B domains have from 10 to 17 well-distributed changes compared to C'.

Structurally, a core of seven critical residues can be identified. These residues are those largely buried upon complex formation or those involved in hydrogen bonds: Glu-35 to Lys-40 from strand β 2 and Tyr-53 from helix α . These seven residues are strictly conserved in eight out of the ten PpL domains (Figure 2). Domains C1 and B5 each have only one nonconservative difference. The changes from Thr-38 to Asn in C1 and from Glu-38 to Thr in B5 could weaken the interaction but not disrupt it. Residue changes outside the structural core in domains C2 and C3 could result in a slightly lower affinity for κ chains compared to that of domain C'. Residues 49 and 52 from domain C' are, respectively, Glu and Arg, which make an internal salt bridge in the middle of the α helix. In domains C2 and C3, these residues are replaced, respectively, by Lys and Ala. This disrupts the salt bridge and places the Lys in close proximity to two Arg residues, and this further results in an unfavorable accumulation of positive charges.

As is the case for the five SpA domains with respect to their interaction with V_L region [13], most PpL domains should also conserve all their hydrogen bonding interactions. Thus, we would expect that all the PpL domains would bind to the V_L region in a similar way so that a whole PpL would contain at least four Ig binding sites. The equivalence in binding between the domains is consistent with previous studies on the binding properties of different PpL constructs. These studies show that avidity increases with the number of Ig binding domains [8]. In fact, a four-domain PpL has an avidity 100-fold higher than the single domain, and a fifth Ig binding domain does not improve avidity further [6]. By superimposing a PpL-Fab complex on each V_L region of whole IgG [27, 28], we can show that the distance between the C terminus of a PpL domain bound to one Fab and the N terminus of the other bound PpL ranges from 60–120 Å depending on the Fab hinge angle. Four domains (including their interdomain linkers of 18 amino

Table 4. Sequences of the Human V_L Gene Families at the Positions Involved in the First Interface with PpL

V _L Subgroup	PpL Binding	L5	L6	L7	L8	L9	L10	L11	L12	L13	L18	L20	L22	L24
κ 1 (19)	+	T	Q	S	P	S	S	L	S	A	R	T	T	R
κ 3 (7)	+	T	Q	S	P	D	S	L	S	LV	R	T	S	R,G
κ 4 (1)	+	T	Q	S	P	D	S	L	A	V	R	T	N	K
κ 2 (9)	–	T	Q	S	P	L	S	L	P,S	V	P	S	S	R,K
λ 1 (5)	–	T	Q	*	P	P	S	V,A	S	G,A,E	R,K	T	S	S,T
λ 2 (5)	–	T	Q	*	P	P,R,A	S	V,A	S	G	S	T	S	T
λ 3 (9)	–	T	Q	*	P	P	S,A	V	S	V	T,M	R	T	S,G,Q
λ 4 (3)	–	T	Q	*	P,S	P,S	S	A	S	A	S	K	T	T
λ 5 (3)	–	T	Q	*	P	P,A,S	S	L,S,H	S	A	S	R,S	T	T,M

The numbers in parentheses for each V_L group indicate the numbers of human germ-line sequences reported in the V Base database (Medical Research Council, Centre for Protein Engineering; <http://www.mrc-cpe.cam.ac.uk/mrc-cpe/>). Families that represent more than 10% of the total genes are reported. Amino acids observed in fewer than 15% of functional germline human genes are not reported. Positions 6 and 7 are shown in *italics*, although they are not in direct contact with PpL to aid the structural alignment between κ 1 and λ chains. The asterisks (*) indicate missing residues. The PpL binding properties are taken from Nilsson et al. [8].

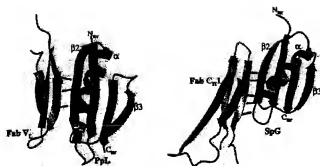


Figure 3. Comparison of SpG-C α 1 and PpL-V α interactions.

Orienting the SpG and PpL β sheets in an analogous manner underscores the structural similarities of both interactions. Strand β 2 of both domains interacts with, respectively, an Fab V α (blue) or an Fab C α 1 (green) β strand (colored lines represent hydrogen bonds between main-chain atoms). In both cases, the bacteriophage strand β 2 forms a continuous β sheet with the Fab through a β zipper interaction.

acids each) could span this distance, and this could explain the strong avidity effect observed on Ig binding. The same pattern of modular domains binding to Igs is shared by SpA and SpG, which like PpL seem to have evolved high avidity for Ig by cassette duplication to allow multiple attachments [29].

Although the K d of the second interface may be substantially higher than the first, we cannot exclude the possibility that the second interface could be structurally important for PpL avidity. A PpL molecule bound to Ig through the high-affinity site of one domain could use the low-affinity site from another domain to interact with the Ig molecules when this would be sterically more favorable than interacting via the first interface.

Structural Convergence between the Fab Binding Modes of SpG and PpL

Streptococcal and *Peptostreptococcal* are found in the same habitats, including the human intestinal and genital tracts. Their cell surface proteins share many common features, and it has been proposed that a transfer of gene fragments may have occurred between these two infective bacteria [29]. With only 15% sequence identity, the Ig binding domains are the least homologous regions between these cell surface proteins. Despite the low sequence identity, these domains share a common fold, with an α helix packed against a four-stranded β sheet. The NMR and crystal structures of both domains have shown that the main structural difference is the orientation of the α helices [7, 30]. The helix runs almost parallel to the β strand direction in the PpL β sheet, whereas in SpG it runs diagonally across the sheet. This arises from a difference in the loop between the α helix and the strand β 3, which is one residue shorter in PpL. Since this is the region of SpG that binds to Fc, it may in part explain the absence of Fc binding by PpL.

The interaction of SpG with C α 1 and that of the first interface of PpL with V α regions of Fab (Figure 3) share similar features. Both domains bury equivalent surface areas upon binding to the Fab and have two common structural features. Firstly, the same region of these two domains is buried in the interactions since SpG also interacts through its strand β 2 [31]. Secondly, in both cases this strand extends a β sheet of the Fab through a β zipper interaction. This interaction involves five main-

chain/main-chain hydrogen bonds in the SpG-C α 1 interface [30] but only three hydrogen bonds in the PpL-V α first interface. The external strand A of the V α region involved in the interaction presents a bulge due to proline at position L8 in the middle of the strand, and this bulge shortens the β zipper. Although the SpG and PpL domains are clearly the result of divergent evolution [29], they have maintained a common binding strategy for interacting with the β strands from different Ig domains (Figure 3) through a main-chain-to-main-chain hydrogen bonding network.

Like SpA, PpL domains target a wide repertoire of human immunoglobulins. As none of those bacterial domains interferes with antibody-antigen recognition, they

Table 6. Summary of Crystallographic Analysis

Data collection ¹	
Resolution (Å)	20–2.6
Unique reflections	29,664
Redundancy	4.66 (5.0)
<i>R</i> _{int}	11.4 (2.6)
<i>R</i> _{meas} (%)	11.8 (46.6)
Completeness (%)	95.2 (93.7)
Space group	P2 ₁ 2 ₁ 2 ₁
Unit cell parameters	<i>a</i> = 55.2 Å, <i>b</i> = 87.3 Å, <i>c</i> = 210.5 Å
Molecular replacement	
Resolution (Å)	20–4.0
Correlation coefficient (%)	91.4 (46.9) ²
Refinement	
Resolution (Å)	20–2.7
<i>R</i> _{meas} (%)	21.5
<i>R</i> _{meas} (%) ³	27.9
<i>R</i> _{int}	
Bonds (Å)	0.007
Angles (°)	1.424
Averaged <i>B</i> factor (Å ²)	45.5
Luzzati coordinates error (Å)	0.34
Number of water molecules	112

*R*_{meas} and *R*_{meas} values in the last shell (2.75–2.7 Å) are, respectively, 31.3% and 34.7%.

²Numbers in bracket give values for the highest resolution shell (2.65–2.6 Å).

³Correlation coefficient for the next highest resolution is indicated in bracket.

⁴Value of *R*_{meas} calculated for 5% randomly chosen reflections not included in the refinement.

may be used to aid the crystallization of antigen-Fab [32, 33] or membrane protein-Fv complexes [34]. We suggest that these small protein domains should be used in a combinatorial manner to increase the variety of possible crystal contacts that can be formed. This would help to solve some problematic crystallization cases. Knowledge of the structure of the PpL-Fab complex will facilitate the engineering of PpL variants with extended binding capacity, for example to encompass λ chains and improve their utility as reagents for Ig detection and purification.

Biological Implications

Several pathogenic bacteria present cell surface proteins with immunoglobulin binding properties that confer on them an advantage during invasion and colonization of host tissues. Some of these bacterial proteins interact with a wide proportion of the immunoglobulin repertoire through interactions with constant regions of immunoglobulin that are highly conserved in sequence. Two bacterial proteins, *Staphylococcus aureus* protein A (SpA) and *Peptostreptococcus magnus* protein L (PpL), interact with a wide range of antibodies by targeting variable, rather than constant, regions of Fab.

We recently reported the crystal structure of an SpA domain in contact with the variable-heavy (V_H) region of an Fab [13] and we now report the crystal structure of a PpL domain in complex with the variable-light (V_L) region of the same Fab. Both bacterial domains interact with the framework part of these variable regions without contacting the hypervariable loops. The positions identified in both complexes account well for the wide repertoire of immunoglobulins recognized by both bacterial domains. In contrast with SpA, which targets conserved residues of the V_H external β sheet, PpL targets a portion of the V_L region that is not strictly conserved in sequence but that maintains a constant backbone conformation among most V_L regions encoded by α genes.

A single domain of this multidomain protein manifests PpL activity. The PpL domain appears to have two independent binding sites for V_L , which interact with very similar areas on the light chain but with markedly different affinities. The residues involved in the V_L interaction are well conserved among the different PpL domains. The interaction of the single domain accounts well for two of the properties reported for the whole protein L. Firstly, PpL exhibits an avidity effect for whole immunoglobulin, and the overall position of a single bacterial domain suggests that a whole PpL with up to five domains in tandem should be able to bind the two V_L regions of an IgG. Secondly, PpL could induce histamine release by bridging two IgE bound to their Fc ϵ receptors at the surface of human basophils or mast cells. The conservation of PpL positions in all domains is also in agreement with this last property.

Experimental Procedures

Crystallization and Data Collection

The 2A2 IgM rheumatoid factor was secreted by a hybridoma line created from myeloma B cells of a rheumatoid arthritis patient [35]. Fab 2A2 was produced by trypsin cleavage of the IgM [35]. The V_L chain of 2A2 belongs to the $\kappa 1$ subtype recognized by PpL. The

PpL domain C' used in the crystallization has 84% sequence identity with the C4 domain from PpL (4 mutations over 51 amino acids) and was produced by recombinant expression [35].

Crystals were grown by vapor diffusion at room temperature in sitting drops by the mixture of a reservoir solution (13%–18% m/v/v/v monomethyl polyethylene glycol (MPEG) 5000, 100 mM imidazole maleate [pH 8.5]) with an equal volume of protein solution (5 mg/ml) containing different Fab 2A2:PpL molar ratios. The ratio was optimized with the aid of streak seeding [37]. Interestingly, the complex crystals grew at a pH that was at or above the upper limit for the growth of unliganded Fab 2A2 crystals. This is different from the Fab 2A2-SpA complex crystallization. In which crystals grew under conditions that were nearly identical to those for the unbound Fab 2A2. Crystals for data collection were enlarged by macroseeding [37].

For data collection, crystals were transferred into a solution containing 50% reservoir solution and 50% cryo-solution (25% m/v/v/v MPEG 5000, 25% v/v/v ethylene glycol, 9% v/v/v xylitol, 100 mM HEPES, pH 7.4). Data were recorded at 120 K from a single crystal on DW32 beamline at LURE synchrotron facility and processed using the HKL package [38]. The crystal belongs to the orthorhombic space group P2₁2₁2, with $a = 55.2$ Å, $b = 87.3$ Å and $c = 210.5$ Å. Statistics are summarized in Table 5.

Structure Determination

We used the molecular replacement method to solve the crystal structure with the program AMoRe [39] by using coordinates of Fab 2A2 (PDB code 1D6B) [13]. The variable (N_L - V_L) and the constant (C_L - C_L) regions were used separately as search models to locate the corresponding parts of the Fab 2A2-PpL complex (Table 5).

After the initial refinement of this model, σ_A -weighted $2F_o - F_o$ and $F_o - F_c$ electron density maps were examined with the program O [40]. The difference maps show electron density for a single PpL domain sandwiched between two Fab fragments. Both the NMR structure of the B1 domain [7] and the crystal structure of the C' domain of PpL (B.J.S., unpublished) were fitted globally in the electron density maps with the best fit for the crystal structure. The 2 Fab:1 PpL model was refined within the 20–2.7 Å resolution range with the program CNS [41] and was rebuilt in O [40]. Detectable PpL domain C' residues range from Glu-20 to Ala-40 (see Figure 2 for numbering). As already observed by NMR [7], the 13 amino acid long N terminus is disordered and thus absent from the final structure. Small loops from the C_L region (H136 to H143 and H186 to H209) of both Fab molecules were poorly defined in maps and removed from the final structure. However, the V_L and V_H regions of both Fabs, the PpL domain, and in particular the V_L -PpL domain interfaces are well defined in the electron density maps (Figure 1b). Statistics for the structure determination are summarized in Table 5.

Mutagenesis

The Y64W PpL and the Y33F-Y64W PpL mutants were constructed with the primers and methods described previously [21]. The D55A-Y64W mutant was made by the mutation of residue 55 on the Y64W template DNA by the use of the antisense primer 5' TGC TAA TAA AGC TGC ATA TOT 3'. The site of the mutation is shown in bold. All mutants were confirmed by DNA sequencing with the Sequencer 2.2 DNA sequencing kit (United States Biochemical Corporation). K_d values determined at pH 8 were obtained by stopped-flow fluorescence measurements as previously described [14].

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Accession Numbers

Coordinates and structure factors have been deposited in the Protein Data Bank (accession code: 1HEZ).

Note Added In Proof

After we submitted this paper, Ulf SJöbring informed us that his laboratory conducted extensive site-directed mutagenesis to document PpL domain folding in collaboration with D. Baker. These experiments, in good agreement with our X-ray structure, gave strong indications that there are two binding sites in protein L: one in the β 2-alpha helix and another one in the β 3 alpha helix (facing the other way).

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Protein and host factors implicated in the pathogenesis of light chain amyloidosis (AL amyloidosis)

Alan Solomon*, Deborah T. Weiss*

* Human Immunology & Cancer Program, Department of Medicine, University of Tennessee Medical Center, Graduate School of Medicine, Knoxville, TN, USA

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Review

Protein and host factors implicated in the pathogenesis of light chain amyloidosis (AL amyloidosis)

Alan Solomon and Deborah T. Weiss

Human Immunology & Cancer Program, Department of Medicine, University of Tennessee Medical Center, Graduate School of Medicine, Knoxville, TN, USA

KEY WORDS: AMYLOIDOSIS, BENCE JONES PROTEINS, PLASMA CELL DYSCRASIA, IMMUNOGLOBULINS, LIGHT CHAINS

ABBREVIATIONS: AL, LIGHT CHAIN-ASSOCIATED AMYLOIDOSIS; I_G, IMMUNOGLOBULIN; V_L, LIGHT CHAIN VARIABLE REGION; C_L, LIGHT CHAIN CONSTANT REGION; J, JOINING; CDR, COMPLEMENTARITY DETERMINING REGION; FR, FRAMEWORK REGION; GDNHCL, GUANIDINE HYDROCHLORIDE; IAPP, ISLET AMYLOID POLYPEPTIDE; PrP, PRION PROTEIN; AEF, AMYLOID ENHANCING FACTOR; IDOR, 4'-ODD-4'-DEOXYDOXORIBUCIN; GAG, GLYCOSAMINOGLYCAN; APO E, APOLOPROTEIN E

Introduction

Light chain-associated (AL) amyloidosis is characterized by the deposition in tissue of monoclonal light chain-related components – a pathologic process that leads to organ failure and eventual death. The lack of information on the etiology of this disease and the fact that there is no effective treatment to prevent or remove the abnormal tissue deposits represent major scientific challenges, the solutions of which may be relevant to other amyloid-associated disorders.

The purpose of this review is to summarize current research efforts that are directed towards elucidation of the protein and host factors thought to be responsible for the induction and perpetuation of AL amyloidosis. The insights that have emanated from these studies and the availability of new technologies, especially in structural and molecular biology, form the basis of future research. The ultimate goal is to apply this information clinically through design of novel therapeutic strategies that will be used to overcome the devastating impact of this disease.

Protein factors in the pathogenesis of AL amyloidosis

Over 20 years have passed since Osserman and colleagues documented the almost invariable presence of serum or urinary monoclonal immunoglobulins (Igs) in patients diagnosed with what was previously termed primary amyloidosis^{1,2}. The prediction that these molecules were involved in the pathogenesis of this disease¹ was confirmed when Glenner and co-workers (and subsequently, many other investigators) demonstrated that the amyloid deposits occurring in such patients were fibrillar in nature and composed of monoclonal light chains or, more commonly, light chain variable-region (V_L) fragments³. The unequivocal relationship between secreted and deposited light chains has since been established through comparative sequence analyses of Bence Jones proteins and amyloid proteins obtained from individual patients^{4,5}.

Historically, research on AL amyloidosis has been directed mainly towards characterizing the primary structural features of monoclonal light chain amyloid-associated proteins – i.e., those extracted from amyloid deposits or excreted in the form of Bence Jones protein, as

Correspondence: Dr. Alan Solomon, University of Tennessee Medical Center, 1924 Alcon Highway, Knoxville, TN 37920, USA
Tel: 423-544-9165; Fax: 423-544-6865; e-mail: asolomon@wizard.hosp.utk.edu

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well as molecules encoded by DNA cloned from bone marrow-derived monoclonal plasma-cell populations. The goal of these studies has been to determine if particular chemical features of the light polypeptide chain render it amyloidogenic. A voluminous body of sequence data has been generated that describes apparently novel amino acid substitutions in the V_L of such light chains⁵. Additionally, in some cases, post-translational modification (e.g., glycosylation) has been thought to account for this phenomenon⁶. However, given the rarity of carbohydrate-containing light chains and the fact that these components do not invariably form amyloid deposits, the functional or pathophysiological effects of glycosylation or other post-translational modifications (deamidation, etc.) are not known.

Light chain primary structure

In an effort to differentiate amyloid proteins from those considered non-amyloidogenic, the primary structures of representative components have been compared. Unfortunately, these studies are confounded by the fact that while there is a relatively large referenced database on the former, the number of documented benign, "non-amyloid," or non-nephrotoxic light chains (Bence Jones proteins that are not associated with myeloma [cast] nephropathy and light chain deposition disease) is markedly limited. Further, those proteins designated "non-amyloidogenic" may, in fact, represent amyloid-associated light chains derived from patients with unrecognized disease.

Another factor that complicates comparative analyses results from the variability inherent in light chain structure:

First, there are two types of light chains, κ and λ , each having distinctive V_L (and constant [C_L]) domains^{5,7}. The V_L portion of the molecule is the product of two exons – V and joining (J) – that encode, respectively, the first ~95 and following ~13 residues. The V_L and V_L domains are characterized by three hypervariable or complementarity-determining regions (CDRs) and four framework regions (FRs) that are involved, respectively, in the antigen-binding site and in maintaining the structural integrity of the molecule (Figure 1). Diversity in sequence results from the presence of multiple, functional V_L and V_L (~35 each) and J_L and J_L (5 and 4, respectively) germ-line genes, as well as the recombinatorial process that links the V and J segments (variation in the length of CDR3 at the V-J joint can also result from the presence at position 96 of non-template-encoded, extra residues that are the products of N or P nucleotides)⁸⁻¹².

Allelic differences can also account for diversity in primary structure, but of greater importance is the pronounced variability that is introduced in light chain-encoded germline sequences by somatic mutation. As yet, there has been no report of an amyloid associated light chain that is completely identical in sequence to that encoded by a V_L or V_L germ-line gene; however, the contributory role of somatic mutation to light chain amyloid formation remains to be established.

Structural features related to amyloidogenicity

What are the structural features that account for the amyloidogenicity of certain light chains? It is noteworthy that in patients with AL amyloidosis λ -type monoclonal

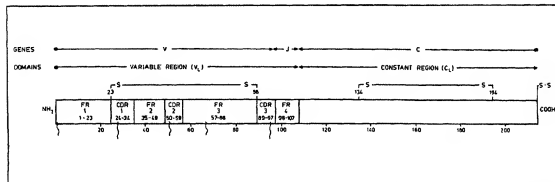


FIGURE 1. Schematic representation of V_L and C_L domains and their genetically-encoded segments. The V_L is the product of two gene segments, V and J, that encode, respectively, the first ~95 residues and the remaining ~13 residues of the V region. The C_L is the product of a single gene, designated C, that encodes for the remaining ~106 residues of the light chain. The location of the three hypervariable regions or CDRs and of the four FRs are as indicated; the J-encoded portion of the V_L encompasses the terminal residue(s) of CDR3 and all the FR4 residues. The wavy lines symbolize the location of additional amino acid residues found within the FRs and CDRs. The residue numbering system is according to Reference 5.

Ig κ predominate, in contrast to the prevalence of κ -type components in other plasma-cell dyscrasias such as multiple myeloma, light chain deposition disease and Waldenström's macroglobulinemia¹⁴. Such overrepresentation of λ chains may be attributed to the fact that these components typically exist as covalent dimers, in contrast to κ molecules that occur in the monomeric as well as dimeric form¹⁵. This difference in quaternary structure may reduce the renal catabolism of λ -type components or, alternatively, increase their propensity to form amyloid. Most remarkable has been the finding that proteins of one particular $V\lambda$ subgroup – $V_{\lambda 101}$ – are invariably associated with this process¹⁶. Despite the relative rarity of this gene family in the normal λ -chain population (~5%), we have documented that 37% (13 of 35) of λ -type Igs obtained from patients with proven amyloidosis were of this isotype¹⁷. Among the primary structural features that distinguish these proteins are the presence of a unique two-residue insertion in FR3 that includes an Asp residue at position 66, as well as a Lys residue at position 17 in FR1. Although the tertiary structural features of $\lambda V1$ light chains have not as yet been elucidated, the interactive potential of these two oppositely charged amino acids may so alter the conformation of these molecules as to render them especially amyloidogenic (F. J. Stevens, personal communication).

To date, comparison of sequence data derived from amyloid and "non-amyloid" proteins (both λ - and κ -type) has failed to reveal a single, site-specific residue that differentiates between these two light chain populations. This finding (as well as the apparent irrelevance of glycosylation) differs from the case of the hemoglobinopathies where, in the case of hemoglobin S, for example, the one residue Glu/Val substitution in the β chain so alters the molecule that when deoxygenated sickling occurs¹⁸.

In contrast to the theory that a single amino acid can render a light chain amyloidogenic, Stevens et al.¹⁹ proposed an alternative mechanism to account for this phenomenon; namely, this process is based on a series of interactions between V_L -associated amino acid residues, first between those in monomers leading to the formation of dimers, then between dimers to form pre-amyloid filaments and finally, between filaments to form fibrils (Figure 2). Because multiple positions within the V_L participate in β -domain interactions that result in V_L -dimer formation and ultimately amyloid fibrils (e.g., salt bridges, hydrogen bonds, or van der Waals contacts), almost any amino acid substitution could theoretically lead to an increase or decrease in the propensity of such interactions to occur, thereby modifying the tertiary structural features of the light chain and causing it, under appropriate physiological conditions, to become amyloidogenic. Conversely, particular amino acid

substitutions could, in fact, inhibit light chain interactions and thus render the protein "non-amyloidogenic." The model satisfies several experimentally defined features of AL amyloid, including fibril diameter and the presence of crossed β sheets that are aligned with the fibril axis and create an intrinsically self-stabilizing assembly.

This hypothesis was supported by the finding that amyloid-associated light chains contained potentially interactive, chemically distinct residues located at key positions within the V_L domain¹⁹. A position-by-position comparison of the frequency of particular residues in 180 human monoclonal κ - and λ -type proteins, 52 of which were obtained from patients with known amyloidosis, revealed statistically significant differences in the CDRs and FRs of the two populations. For example, amyloid-associated molecules were most often characterized by the presence in the CDRs of sterically accessible, charged residues (e.g., Asp). Alternatively, in FR2, the typical basic residue found at position 45 in non-amyloid proteins was replaced by Asn and in FR1 at position 20, amyloid constituents had a hydrophobic Ile residue instead of Thr or Ser. For both κ and λ light chains, the majority of amyloid-associated positions were distributed along the surface of the light chain involved in the antigen-binding site and, therefore, would not participate in the internal packing of the V_L domain. Further, the side chains of residues at such positions would extend into the solvent and be accessible for interaction with other molecules in solution.

These chemical differences that potentially modify light chain tertiary structure were also predicted to enhance V_L -dimer formation and result in protein aggregation. Earlier studies, using size-exclusion chromatography, demonstrated that, on the basis of polymer formation, Bence Jones proteins obtained from patients with amyloidosis or other types of pathologic light chain deposits could be distinguished from non-toxic components²¹. The demonstration that amyloid-associated proteins share unique primary structural features and readily aggregate in an *in vitro* system provides further evidence for the three-step molecular mechanism by which light chains form fibrils, i.e., dimer \rightarrow filament \rightarrow fibril^{19,22}.

Wetzel and colleagues²³ have hypothesized a somewhat different theory to explain light chain amyloidogenesis. They posited that, while multiple substitutions at key sites in the V_L are responsible for light chain amyloidogenicity, this phenomenon of amyloid fibril formation results because such replacements so modify the tertiary structure that the protein becomes partially or completely unfolded. It is this intermediate form, then, that is the culprit in AL amyloid formation (as has been postulated to occur in the transthyretin-associated amyloidosis²³). Experimental support for this theory was obtained using V_L fragments

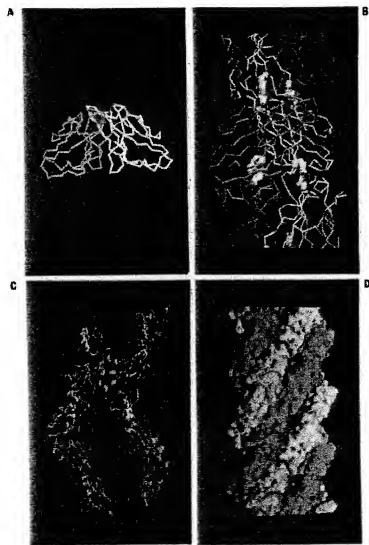


FIGURE 2. Computer-based models of V_L interactions leading to the formation of amyloid fibril. (A) Backbone representation of a V_L dimer. This basic structure has been found for most Ig light chains which have been studied crystallographically. The shading surrounding the "backbone" indicates the volume occupied by atoms. The portion of the molecule that constitutes the equivalent of an antibody's antigen-binding site is located at the "top" of the molecule in this representation; among light chains, most amino acid substitutions occur in this area. (B) Hypothetical arrangement of V_L dimers during the formation of non-covalent polymers. The antigen-binding site of one dimer interacts with the surface of the axially-opposed end of a second dimer following a 90° rotation about the two-fold symmetry axis. Many amyloid-forming light chains have amino acids of opposite charge, as highlighted in this figure, located in positions to form high-affinity "salt bridges." A consequence of the linking of V_L domains is that any weak "non-specific" affinity of the dimer surface for particular tissue surface features, such as proteoglycans, is amplified. Thus, a light chain, which individually would not interact with an organ or tissue, might be specifically bound to an organ when polymerized. (C) Interaction of V_L -dimer filaments. When viewed from a different angle, the filament shown in (B) has a convoluted or "sawtooth" surface that makes it possible to bring together two or more filaments. In this figure, an anti-parallel mechanism of bringing filaments together is illustrated; several amino acid positions at which substitutions could either enhance or suppress the docking of filaments are highlighted. (D) Space-filling representation of the proposed V_L -dimer polymerization model. (Figure furnished by Dr. Fred J. Stevens, unpublished data.)

derived from DNA recombinant technology. These molecules, expressed in a bacterial system, were based on the V_L dimer REI – a presumably non-amyloidogenic protein, the tertiary structure of which had been elucidated by x-ray crystallography²⁴. Six V_L constructs were prepared that contained replacements thought to be unique in κ - or λ -type amyloid-associated light chains. These modifications, due to their locations, were expected to destabilize the folded structure of the V_L domain. In contrast to the wild-type REI component, all six mutants, at low pH, formed aggregates and bound Congo red. When examined in the presence of guanidine hydrochloride (GdnHCl), four of the six were unstable and aggregated to produce amyloid-like fibrils. The most destabilizing effect occurred when a highly-conserved Arg at position 61 was replaced by Asp. This substitution was predicted to eliminate or decrease the potential to form a salt bridge/hydrogen bond with another Asp residue located at position 82 in an adjacent loop. Although the GdnHCl *in vitro* stability assay may provide a means to predict the fibrillogenic capacity of a Bence Jones protein, the relevance of this system to *in vivo* amyloid formation remains to be established.

The aforementioned studies involving comparative analyses of light chain sequence have made it increasingly evident that such data *per se* are not sufficient to differentiate amyloid from non-amyloid components. In order to define more precisely the relationship between particular amino acid residues and light chain amyloidogenicity, it is necessary to determine how variations in sequence affect the overall tertiary structure of the molecule.

Because amyloid proteins cannot be crystallized, computer modeling offers a powerful new tool to predict the three-dimensional impact imparted on a protein by modification of and interactions between amino acids. Additionally, the ability to generate *via* recombinant technology a replenishable source of human light chains in which primary structural modification can be induced represents an important means to advance our understanding of the protein basis of amyloidogenicity^{22,23,24}. This technique, as well as computer technology, provides exciting new approaches to research in this field, since manipulation of key amino acids and, most importantly, visualization of resulting interactions should provide information on why certain proteins form amyloid.

Light chain fragmentation in AL amyloidosis

The structural requirements for the *in vivo* conversion of soluble light chain precursor proteins into insoluble amyloid fibrils have not been definitely established – e.g., it is unknown if the native protein itself can form amyloid fibrils *in vivo* or if this process is initiated or accelerated by light chain fragmentation. In rare instances, the protein

extracted from AL amyloid deposits was found to be composed exclusively of an intact light chain^{2,27}. More commonly, these extracts consist of light chain fragments (V_L or V_L plus a portion of the C_L). It has not been conclusively established if these components are produced *de novo*^{24,29} or result from catabolism or *in situ* proteolysis of intact light polypeptide chains. The fact that we have found in virtually all AL amyloid extracts examined a certain, albeit small, amount of the complete light chain suggests that the common occurrence of fragments represents a degradative process. It is noteworthy that, based on the tertiary structural features of light chains, as evidenced by x-ray crystallography, the C-terminal residues of amyloid fragments (in positions 152-154) are typically located in sterically exposed regions of the molecule; such sites would thus be accessible for endopeptidase digestion. We have been able to generate similarly sized fragments using various types of endopeptidases (e.g., cathepsin D, pepsin, or that contained in a lysosomal extract prepared from kidney). Additionally, we have found that amyloid fragments isolated from different organs from the same patient can vary in molecular mass³⁰. Although such variations may reflect deposition of synthetically-derived light chain fragments, it is also possible that light chains are deposited in a relatively intact form as amyloid and that degradation occurs as mediated by local tissue factors. If light chain proteolysis is, indeed, essential for amyloid formation, the presence or extent of light chain fragmentation could have prognostic or therapeutic implications.

Experimentally, it has been shown that endoprotease digestion of certain Bence Jones proteins obtained from patients with or without amyloidosis yielded V_L fragments that had the characteristic features of amyloid^{31,34}. This finding may indicate that the C_L portion of the molecule interferes with amyloid formation or, as has been shown *in vitro*, is remarkably susceptible to proteolysis as compared to the V_L domain³⁵. The demonstration that regions within the light chain are potentially amyloidogenic has come from studies of Eulitz and co-workers³⁶⁻³⁸, who found that tryptic digestion of certain extensively reduced and alkylated amyloid-associated Bence Jones proteins yielded precipitates that exhibited green birefringence after Congo red-staining and were fibrillar when viewed by polarization light microscopy and electron microscopy, respectively. The insoluble tryptic digests, when rendered soluble in trifluoroacetic acid and subjected to HPLC-chromatographic separation on a reversed-phase column, were found to contain characteristic 18- to 43-residue V_L - and C_L -related peptides (this discovery was analogous to the observation that synthetic 10- to 28-residue peptides corresponding to portions of the Alzheimer's disease

amyloid-associated β protein, the islet amyloid polypeptide [IAPP], or prion protein (PrP) could form amyloid fibrils *in vitro*³⁹⁻⁴¹). More recent studies in our laboratory have led to the identification and synthesis of 12- to 24-mer V_L -related peptides that are particularly fibrillogenic (Ref. 30 and Niewold, Th. A. et al., unpublished studies). These findings suggest that, under appropriate conditions, all light chains are potentially amyloidogenic and that the susceptibility of these components to proteolytic cleavage may depend on other factors involved in light chain catabolism. That certain Bence Jones proteins can themselves function as peptidases⁴⁴ could also be of pathologic significance; however, the relevance of this phenomenon to the *in vivo* production of amyloid remains to be established.

If, indeed, 12- to 24-mer "amyloid" peptides are generated *in vivo* through proteolysis of precursor monoclonal light chains, such components may contribute to the pathogenesis of AL amyloidosis. Lansbury and colleagues⁴⁵⁻⁴⁸ have proposed that amyloid formation requires or is accelerated by a seeding mechanism whereby preformed amyloid fibrils serve as a nucleation substance that initiates conversion of a soluble precursor protein or peptide into amyloid. Three factors were deemed essential for a nucleation-dependent polymerization process to occur: First, a lag time is necessary before detectable aggregates appear; second, there must be a critical concentration of the monomer before polymerization occurs; and third, polymerization is accelerated by a preformed "seed" or nucleus. That this phenomenon may ensue *in vivo* has been evidenced by the rapid formation of amyloid in the experimental hamster and mouse AA models^{49,50} through the administration of "amyloid-enhancing factor" (AEF) - a sonicated preparation of an organ that contains either AA- or AL-related amyloid fibrils.

However, *in vitro* experiments have indicated that seeding is protein-specific; i.e., there must be complementarity between seed and some portion of the precursor protein for growth to occur, as shown in the case of BA4 and prion fibrillogenesis^{45,51-53}. The demonstration that amyloid formation can be accelerated by an apparent nucleation-dependent polymerization mechanism provides, as suggested by Jarrett and Lansbury⁴, a rational therapeutic approach - namely, reduction of precursor protein concentration and interference with the generation of the nucleus should slow this relentless process. In this regard, α_1 -antichymotrypsin and a silicate compound (Na_2SiO_3) were found to inhibit BA4 fibrillogenesis^{54,55}. Another agent - 4'-iodo-4'-deoxydoxorubicin (IDOX), an iodinated anthracycline compound was shown to bind with high affinity to AL as well as other types of amyloid deposits and also interfered with *in vitro* fibrillogenesis. When

incubated with AEF, this agent significantly reduced AA amyloid formation in an *in vivo* experimental animal system⁵⁶. Further, five of eight patients with AL amyloidosis who were treated with this compound had objective evidence of amyloid resorption, despite no diminution in Bence Jones proteinuria⁵⁷.

Pathologic heterogeneity of light chain deposits

Although the three major forms of the human light chain-associated diseases (myeloma [cast] nephropathy, light chain deposition disease and AL amyloidosis) exist as discrete entities^{14,15,39}, the coexistence of two distinct forms of light chain-associated disease - e.g., light-chain deposition disease and amyloidosis - has been noted^{19,42}. It is not known whether each form results from the presence in individual patients of multiple Bence Jones protein components, from the potential of a single Bence Jones protein to induce more than one type of pathology, or from a natural progression of one disease state to another, as evidenced by the common involvement of blood vessel walls in non-fibrillar and fibrillar forms of light chain deposition.

The fact that a monoclonal light chain can assume multiple conformers according to the solvent used for crystallization^{19,61} provides evidence that physiological factors can profoundly influence V_L dimer interaction. The failure of a particular Bence Jones protein to form amyloid might be attributed to the greater propensity of the protein to aggregate as amorphous casts (myeloma [cast] nephropathy) or as punctate, linear deposits (light chain deposition disease). Alternatively (as previously discussed), particular amino acid substitutions may prevent V_L aggregation or, if light chain fragmentation is a prerequisite for amyloid formation, such substitutions may so modify the native conformation of the light chain that protease-sensitive sites in the C_L domain are rendered inaccessible. Furthermore, local factors may change the protein thus deposited: Initially the deposits may be non-fibrillar but are subsequently modified and assume the β -pleated structure typical of amyloid fibrils.

Host factors in the pathogenesis of AL amyloidosis Accessory Molecules

Interactions between monoclonal light chains and other biologically active molecules may also lead to pathologic deposition. For example, myeloma (cast) nephropathy has been thought to be the product of precipitated complexes comprised of Bence Jones protein and a component produced by the renal distal tubules - Tamm-Horsfall protein^{64,66}. In the case of AL amyloid, the deposits, in addition to light chains, have been shown to contain several

chemically diverse substances, including amyloid P component⁴⁷, the highly-sulfated glycosaminoglycans (GAGs) heparan sulfate and dermatan sulfate⁴⁸⁻⁵⁰ and apolipoprotein-E (Apo-E)⁵¹. Other types of molecules, such as ubiquitin, complement-related components, collagen, fibronectin and laminin, have been detected as well. The demonstration that certain of these compounds (e.g., Apo-E) can accelerate fibril formation *in vitro* has led to their designation as "pathologic chaperones"⁷¹⁻⁷³. It has been proposed that these molecules induce a β -pleated-sheet conformation of the precursor protein that, in turn, leads eventually to fibril formation. However, we and others^{33,34,36-38,74} have demonstrated that isolated light chains and related fragments can form amyloid fibrils *in vitro* in the absence of ancillary compounds. Thus, it is possible that the effects of these substances are secondary; i.e., binding is non-specific and results from the highly adsorptive nature of the β -pleated-sheet structure of the amyloid fibril. The "pathologic chaperones," which are not found in non-amyloid forms of light chain deposition^{62,72}, may not necessarily facilitate fibrillogenesis but, rather, participate in other functions that are part of the amyloidogenic process – e.g., GAGs may play an important role in selective organ localization (see below) of AL amyloid deposits and P component has been implicated as a factor that can inhibit or prevent the resolution of this material⁷⁵.

Light chain metabolism

Essential to the pathogenesis of AL amyloidosis is a continuous endogenous source of precursor protein, i.e., the monoclonal light chain. However, the amount of protein required for this process is unknown. Since the catabolic half-life of light chains is normally <2 h, the concentration of these components in serum or urine reflects only a small percentage of the protein synthesized by the monoclonal plasma-cell population⁷⁶. In contrast to patients with multiple myeloma, individuals with AL amyloidosis most often have a low percentage of bone marrow plasma cells (median 5%) and minimal Bence Jones proteinuria (median 0.4 g/24 h)⁷⁷. Although there is evidence to suggest that a population of monoclonal B cells circulates and serves as precursor to the bone marrow-derived plasma cells⁷⁸, it is unlikely that there exists a significant extramedullary cellular source of monoclonal light chains (except in the case of localized AL amyloid deposits⁷⁹). There is as yet no evidence to indicate that the light chain synthetic rate of monoclonal plasma cells obtained from patients with AL amyloidosis is greater than that of plasma cell populations associated with other forms of pathologic light chain deposition. The fact that patients with AL amyloidosis can have extensive disease despite a seemingly low serum or urine concentration of precursor proteins suggests (in the

absence of an inordinately high rate of synthesis) an exceptional propensity of these molecules to form amyloid. It is also possible that such patients may have unusually high concentrations of light chain binding factors⁸⁰⁻⁸² or a molecular defect that limits or prevents the elimination of the AL deposits.

Organ diversity of AL deposits

Central to the pathophysiology of this disease is the remarkable diversity that exists in the organs affected in patients with AL amyloidosis^{77,83}. In some individuals, the deposits are confined principally to the kidney and in others to the heart, small intestine, or peripheral and/or sympathetic nerves, etc. Further, they may be primarily vascular, interstitial, or both. Our analyses of protein extracted from spleen and other organs have shown no relationship between the molecular mass of the deposited material and the affected tissue⁸⁴. Whether selective tissue affinity is related to specific primary structural features of the light chain that result in an interaction with local tissue factors or to an antibody-like affinity of certain light chains for particular tissue constituents remains to be established. Previous attempts to demonstrate organ specificity *in vitro* using fluorescein-labeled native Bence Jones proteins obtained from patients with particular organ involvement, although showing qualitative differences, were considered inconclusive⁸⁵.

The demonstration by x-ray crystallography that Bence Jones proteins and V_L dimers can structurally mimic Fab fragments, including the antigen-binding site⁸⁶, implies that the selective organ deposition of light chains may represent an antibody-ligand interaction. Alternatively, AL amyloid deposition may be due to the synthesis of the amyloidogenic precursor protein by a local monoclonal plasma-cell population. This phenomenon would be analogous to the site-specific production of other types of amyloid – e.g., A₁AI, AANF, AIAPP, A β – associated with precursor proteins (calcitonin, atrial natriuretic factor, islet amyloid polypeptide, BA4 protein, respectively). Another mechanism that may account for the localization of AL amyloid deposits within particular organs involves the interaction between precursor light chains and tissue-specific molecules – e.g., GAGs. It remains to be determined if protein structure, organ affinity, or local production of light chains, etc. is responsible for the clinical diversity found in localized amyloid deposition.

Pathophysiology of AL amyloidosis

Other physiologic factors analogous to those implicated in myeloma (cast) nephropathy may be involved in the pathogenesis of AL amyloidosis. For example, the precipitation of Bence Jones proteins in the form of renal

tubular casts is accelerated by dehydration or other conditions that can adversely affect renal function – e.g., hypercalcemia, anemia, or infection¹⁴. Due to the profound effect of solvent composition on light chain tertiary structure⁶², it is possible that, under appropriate conditions, soluble precursor light chains can be induced to form amyloid fibrils.

Other host (patient)-related elements that could account for AL amyloid pathogenesis include the local processing by macrophages of soluble Bence Jones protein to form amyloid fibrils⁶⁴ or the failure of these cells to remove or prevent fibrillar light chain deposits. In contrast to the typical giant-cell reaction to Bence Jones protein-containing casts in myeloma (cast) nephropathy, characteristically, there is scant inflammatory response to AL amyloid deposits, presumably since this material is weakly or non-immunogenic^{65,66}. With rare exception, AL amyloid deposits are apparently irreversible. The failure of the host to degrade AL amyloid fibrils may result from a deficiency in macrophage function or lack of a requisite proteolytic enzyme (as postulated in the pathogenesis of brain amyloid in Alzheimer's disease^{67,68}). The coating of amyloid fibrils by "pathologic chaperones" – e.g., P component⁶⁹ – may also interfere with fibril degradation. Although there is no documented difference in the association of such components with λ - or κ -type amyloid fibrils, it has been reported that α 1-antitrypsin was capable of disaggregating λ - but not κ -chain-containing fibrils⁶⁹.

In Vivo models of AL amyloidosis

Further insight into the pathogenesis and, ultimately, the effective treatment of AL amyloidosis depends, in part, on the availability of *in vivo* experimental models that duplicate the human form of this disease. We have reported the results obtained with one such model whereby mice were repeatedly injected with different Bence Jones proteins obtained from patients with AL amyloidosis. In these experiments, the human proteins were deposited in the kidney and other organs of the recipient animals in the form of amyloid^{90,91}. The human light chain amyloid also contained mouse amyloid P component. Conversely, no deposits were found in mice injected under similar conditions with a "control," i.e., non-amyloid, Bence Jones protein. Due to the relatively large amount of protein needed for injection and the length of time required for the amyloid to form, other models in which human amyloid-associated light chains are produced transgenically or by transfectomas may prove more suitable. The development of light chain-related amyloid in such *in vivo* models will provide an invaluable means for further research on the pathogenesis, treatment and prevention of AL amyloidosis.

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Partially Folded Intermediates as Critical Precursors of Light Chain Amyloid Fibrils and Amorphous Aggregates[†]

Ritu Khurana, Joel R. Gillespie,[‡] Anupam Talapatra, Lauren J. Minert, Cristian Ionescu-Zanetti, Ian Millett,[‡] and Anthony L. Fink*

Department of Chemistry and Biochemistry, University of California, Santa Cruz, California 95064

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ABSTRACT: Light chain, or AL, amyloidosis is a pathological condition arising from systemic extracellular deposition of monoclonal immunoglobulin light chain variable domains in the form of insoluble amyloid fibrils, especially in the kidneys. Substantial evidence suggests that amyloid fibril formation from native proteins occurs via a conformational change leading to a partially folded intermediate conformation, whose subsequent association is a key step in fibrillation. In the present investigation, we have examined the properties of a recombinant amyloidogenic light chain variable domain, SMA, to determine whether partially folded intermediates can be detected and correlated with aggregation. The results from spectroscopic and hydrodynamic measurements, including far- and near-UV circular dichroism, FTIR, NMR, and intrinsic tryptophan fluorescence and small-angle X-ray scattering, reveal the build-up of two partially folded intermediate conformational states as the pH is decreased (low pH destabilized the protein and accelerated the kinetics of aggregation). A relatively native-like intermediate, *I_N*, was observed between pH 4 and 6, with little loss of secondary structure, but with significant tertiary structure changes and enhanced ANS binding, indicating exposed hydrophobic surfaces. At pH below 3, we observed a relatively unfolded, but compact, intermediate, *I_C*, which was characterized by decreased tertiary and secondary structure. The *I_C* intermediate readily forms amyloid fibrils, whereas *I_N* preferentially leads to amorphous aggregates. Except at pH 2, where negligible amorphous aggregate is formed, the amorphous aggregates formed significantly more rapidly than the fibrils. This is the first indication that different partially folded intermediates may be responsible for different aggregation pathways (amorphous and fibrillar). The data support the hypothesis that amyloid fibril formation involves the ordered self-assembly of partially folded species that are critical soluble precursors of fibrils.

Immunoglobulin (Ig)¹ light chains are involved in several protein deposition diseases, including one resulting in the formation and deposition of amyloid fibrils (light chain or AL amyloidosis) and another known as light chain deposition disease that involves amorphous protein deposits (*1, 2*). The morphology of the deposited aggregates in these two diseases is clearly different, and typically patients exhibit only one

form of light chain deposition. However, there is at least one report of a patient exhibiting both AL amyloidosis and LCDD involving the same light chain (*3*). The exact length of light chains in amyloid deposits varies, but is usually in the 110–130 residue range (12–14 kDa) corresponding to the intact variable domain (*4*).

The molecular mechanisms leading to amyloid formation are poorly understood. In this report, we address the question of why some immunoglobulin light chains form amyloid and related deposits while others do not, in particular the hypothesis that protein aggregation arises from the self-association of partially folded intermediates. Support for this hypothesis has been found with proteins such as transthyretin (*5*) and lysozyme (*6*). We postulate that amyloid fibril formation from native proteins occurs via a conformational change leading to formation of a partially folded intermediate conformation, association of this intermediate to form soluble oligomers leading to the critical nucleus, and subsequent formation of the initial fibrillar species, typically a filament or protofibril, and finally association of protofibrils into mature fibrils (*7*).

We have investigated the biophysical properties and amyloidogenicity of the variable domain of a recombinant amyloidogenic light chain, SMA, engineered by Stevens et al. (*8*). SMA (114 residues, *M_r* = 12 700) was initially

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* Correspondence should be addressed to this author at the Department of Chemistry and Biochemistry, University of California, Santa Cruz, CA 95064. Tel: (831) 459-2744; Fax: (831) 459-2935; E-mail: enfink@cats.ucsc.edu.

[‡] Current address: Department of Chemistry, Stanford University, Stanford, CA 94305.

¹ Abbreviations: SAXS, small-angle X-ray scattering; *R_g*, radius of gyration; FTIR, Fourier transform infrared spectroscopy; ATR, attenuated total reflectance; IRE, internal reflectance element; TEM, transmission electron microscopy; AFM, atomic force microscopy; TFT, Thioflavin T; ANS, 8-anilino-1-naphthalenesulfonate; CD, circular dichroism; Ig, immunoglobulin; V_L, variable domain of Ig light chain.

extracted from lymph node-derived amyloid fibrils of an AL amyloidosis patient (9). A very similar light chain domain, LEN, was derived from a patient with multiple myeloma who showed no evidence of renal dysfunction or amyloidosis (10). The IgG-V_L domains consist of a highly conserved framework formed by two sheets of antiparallel β -strands forming a β -sandwich, and three loops comprising the complementarity-determining regions (CDR) that form part of the antigen binding site. The sequences of SMA and LEN are very similar, differing only at 8 positions out of 114. Four of these are in CDR3 (Q89H, T94H, Y96Q, S97T), two are in CDR1 (S29N, K30R), and the remaining two are in the framework region (P40L, I106L). The high-resolution crystallographic structure of LEN (1.8 Å) has been solved (PDB Accession No. 1LVE) (11). Both SMA and LEN belong to the κ IV family of Igs.

The amyloidogenic light chain, SMA, is significantly less thermodynamically stable than LEN under all conditions (unpublished observations). The presence of low concentrations of denaturants also results in fibril formation from the "benign" LEN (12). In the present study, we used biophysical characterization of the conformation of SMA as a function of pH to reveal the presence of two distinct partially folded intermediates: one with relatively native-like properties, the other with relatively unfolded properties. Destabilizing conditions at physiological pH, e.g., low urea concentration, also lead to aggregation and fibril formation. Thus, conditions that result in population of these intermediates lead to aggregation, supporting the hypothesis that partially folded intermediates are key precursors on the aggregation pathway. Interestingly, both amorphous and fibrillar aggregates were observed, and were shown to arise from two different intermediates.

MATERIALS AND METHODS

Expression and Purification of Recombinant V_L SMA. The recombinant V_L domain SMA was purified from JM83 *E. coli* cells transformed with the plasmid pKIVsma004, generously provided by Dr. Fred Stevens, Argonne National Lab (8). The plasmid construct was based on the pASK vectors constructed by Skerra et al. (13) and contained an *ompA* leader for periplasmic localization of the protein to ensure the formation of the core disulfide bond. The overexpressed protein was purified using the procedure of Stevens et al. (8) with minor modifications. Briefly, the recombinant protein was extracted from the periplasm using osmotic shock via treatment with ice-cold TES followed by distilled water. The periplasmic extract was dialyzed against 4 changes of 20 volumes of 10 mM acetate buffer, pH 5.6, and loaded onto a fast-flow SP Sepharose column. The column was washed with 10 mM acetate buffer, pH 5.6, and the protein eluted using 10 mM phosphate buffer, pH 8.0. The fractions were assayed by SDS-PAGE, and fractions containing the recombinant protein were pooled, filtered through 0.22 μ m filters, and stored in glass vials. Typical yields were 7–8 mg of purified protein per liter of cells. Protein concentrations were measured via optical density at 280 nm using the extinction coefficient of $E_{0.1\%}^{1\text{cm}} = 1.8$ calculated from the sequence. The purified protein was stored in 10 mM phosphate buffer (pH 8.0) at 4 °C and used within 2 weeks of the initial purification. The purity of the protein prepara-

tions was assayed by SDS-PAGE and by electrospray mass spectrometry (Micromass Quattro II).

Intrinsic Tryptophan Fluorescence Measurements. Fluorescence measurements were made with a FluoroMax-2 fluorescence spectrometer (Jobin Yvon-Spex). Emission spectra between 300 and 420 nm were collected with excitation at 280 nm. Spectra were collected at different pHs within the range of 10–2 using 0.5 μ M protein samples in 50 mM of the appropriate buffer containing 100 mM NaCl. Spectra were collected at 25 and 37 °C. The stability of SMA toward urea denaturation was monitored as a function of pH by recording changes in tryptophan fluorescence intensity upon excitation at 280 nm and emission at 350 nm at 25 °C.

Samples of SMA (1 μ M monomer) were incubated in 20 mM phosphate buffer (pH 7.4), 100 mM NaCl containing varying amounts of urea (0–8 M) for 2 h to ensure completion of the unfolding reaction. Data were analyzed by nonlinear least-squares fitting to a two-state folding model. The fraction unfolded, F_u , was calculated using the equation: $F_u = (y_t - y_f)/(y_t - y_u)$ where y represents the observed fluorescence at a particular concentration of urea, and y_t and y_u represent the corresponding fluorescence of the folded and unfolded states, respectively, at that urea concentration. For baseline fitting, a linear least-squares analysis was performed to determine the values of y_f and y_u as a function of urea concentration. The free energies of unfolding were calculated as a function of urea concentration using the equation: $\Delta G = -RT \ln K_{eq}$, where $K_{eq} = f_u/(1 - f_u)$. ΔG° was determined by linear extrapolation to zero urea concentration using the expression: $\Delta G^\circ = \Delta G + m[\text{urea}]$.

ANS Binding. 1,8-Anilinonaphthalenesulfonate was obtained from Kodak, and a stock solution was prepared by dissolving in water followed by filtration through a 0.2 μ m syringe filter and measuring the concentration using an extinction coefficient of 5000 M⁻¹ cm⁻¹ at 350 nm. The fluorescence emission spectra of solutions of 10 μ M ANS and 0.5 μ M protein were collected between 420 and 600 nm upon excitation at 380 nm as a function of pH at 37 °C.

Circular Dichroism Spectra. CD spectra were collected on an AVIV model 62DS spectrometer between 260 and 190 nm for the far-UV region and between 320 and 250 nm for the near-UV region, with a step size of 0.5 nm and an averaging time of 5 s and collecting 5 repeat scans. Cells of 1 and 0.01 cm path length were used for near- and far-UV CD measurements with protein concentrations of 0.5 and 1.7 mg/mL, respectively.

pH Dependence. pH-dependent changes in spectroscopic data were fit using a modified Henderson-Hasselbalch equation for one (eq 1) or two (eq 2) transitions, to determine the midpoints of the transitions:

$$Y_{\text{obs}} = \frac{Y_N + Y_U 10^{\text{pH} - \text{pH}_{\text{H}_2}}}{1 + 10^{\text{pH} - \text{pH}_{\text{H}_2}}} \quad (1)$$

$$Y_{\text{obs}} = \frac{Y_N + Y_U 10^{\text{pH} - \text{pH}_{\text{H}_1}} + \frac{Y_{\text{IV}}}{10^{\text{pH} - \text{pH}_{\text{H}_2}}}}{1 + 10^{\text{pH} - \text{pH}_{\text{H}_1}} + \frac{1}{10^{\text{pH} - \text{pH}_{\text{H}_2}}}} \quad (2)$$

where Y_{obs} is the observed spectroscopic property, Y_N is the

value of the spectroscopic property for the native state, Y_N , is the spectroscopic property for the native-like intermediate, and Y_U is the spectroscopic property for the unfolded-like intermediate. pH_{m1} and pH_{m2} are the midpoints of transitions from the native state to the I_N intermediate and from I_N to the I_U intermediate, respectively (36).

Thin film ATR-FTIR measurements were performed using a SPECAC out-of-compartment ATR accessory and a Nicolet 800SX FTIR bench. A germanium crystal IRE was used for making hydrated thin films of ~50–100 μ g of protein from both soluble and insoluble protein as previously described (14, 15). ATR-FTIR spectra were collected followed by Fourier transformation of the sample spectra using a clean crystal spectrum as a background. The water vapor spectrum was collected by reducing the air purge and subtracted from the protein spectrum until the spectra were featureless in the region between 1700 and 1800 cm^{-1} . ATR-FTIR spectra for SMA were collected at pH 7.5 in 50 mM sodium phosphate, 100 mM NaCl, at pH 5.0 in 50 mM sodium acetate, 100 mM NaCl, and at pH 2.0 in 20 mM HCl, 100 mM NaCl. Buffer spectra were subtracted from the sample spectra. Component spectra were obtained by first determining peak positions using both second-derivative and Fourier-self-deconvoluted spectra, followed by curve-fitting to the raw spectrum (16).

In Vitro Fibril Formation Assays. Fibril formation was monitored using a fluorescence assay based on the enhanced fluorescence of the dye Thioflavin T (TFT) on binding to amyloid fibrils (17). Amyloid fibrils were grown from purified protein (40 μ M) in 50 mM buffer and 100 mM NaCl. A filtered protein sample (using 0.22 μ m syringe filters) was incubated under the desired conditions in a 1.8 mL flat-bottomed screw-capped glass vial with moderate stirring using a Teflon-coated micro-stir bar. Typical fibril growth experiments involved incubating the protein at 37 °C with constant stirring and removing aliquots (10 μ L) over time for analysis by light scattering and TEM and TFT binding (see below). Standard buffers included 20 mM HCl or phosphate (pH 2), 50 mM formate (pH 3 and 4), 50 mM cacodylate or acetate (pH 5 and 6), and 20 mM TRIS or 20 mM HEPES or 50 mM phosphate (pH 7). Both Rayleigh light scattering and fluorescence spectra were collected using a SPEX/Jobin-Yvon Fluoromax-2 spectrofluorometer. Constant temperatures were maintained using a circulating water bath. At each time point, a 220 μ L sample was removed and transferred to a cylindrical quartz microcell to measure Rayleigh light scattering at 330 nm with a 1 nm band-pass for both excitation and emission monochromators. Thioflavin T binding assays were conducted by adding sample aliquots (10 μ L) to 990 μ L of 20 μ M TFT in 50 mM TRIS, pH 7.5, and 100 mM NaCl in a 1 mL fluorescence cuvette. Fluorescence emission was monitored with excitation at 450 nm using a 5 nm band-pass on both the excitation and emission monochromators. Fluorescence intensities were reported at 482 nm.

SAXS Measurements. Small-angle X-ray scattering measurements were performed on beam line 4-2 at the Stanford Synchrotron Radiation Laboratory (SSRL) as described previously (18). The SAXS instrument was configured with a Mo/CB, multilayer monochromator, an 18 mm beamstop, and a 218 cm sample-to-detector distance. Data were collected on protein samples ranging from 0.5 to 10 mg/mL

in 50 mM buffer and 100 mM NaCl at 37 °C using a PTFE flow-cell with 1.3 mm path length to minimize radiation damage. Radii of gyration were calculated using the Guinier approximation (19).

Atomic Force and Transmission Electron Microscopy. Transmission electron micrographs were collected using a JEOL JEM-100B microscope operating with an accelerator voltage of 80 kV. Typical nominal magnifications ranged from 27000 \times to 67000 \times . Samples were deposited on Formvar-coated 300 mesh copper grids and negatively stained with freshly prepared 2% aqueous uranyl acetate.

For AFM, aliquots of 50 μ L of incubation solution were transferred to an Eppendorf tube and spun to pellet the precipitated material, which was then washed twice with water before resuspending in deionized water. A drop of aggregate suspension was deposited on freshly cleaved mica and dried immediately with nitrogen gas. The samples were imaged with an Autoprobe CP AFM (Park Scientific, Sunnyvale, CA) in the noncontact (NC-AFM) mode. The tube scanner was a 100 μ m Scanmaster (Park Scientific). NC Ultralevers (Park Scientific) were used as cantilevers. The resonant frequency was ~100 kHz. The images were taken in air, ambient conditions, at a scan frequency of 1–2 Hz, using silicon nitride tips.

NMR Spectroscopy. ^1H NMR spectra were collected using a Varian Unity+ 500 spectrometer equipped with ultraslims and a Varian triplet resonance probe. Presaturation and postacquisition digital filtering were used for solvent suppression. Data were collected on 0.5 mL protein samples containing 100 mM NaCl and 10% D₂O. The pH was adjusted by titration with NaOH or HCl as needed. All data were recorded at 37 °C.

Light Scattering Measurements at High SMA Concentration. The rate of aggregation was monitored by static light scattering using a Nepheloskan instrument from Labsystems and a 96-well plate reader. Solutions of 3.5 and 7 mg/mL SMA at the appropriate pH were incubated at 37 °C along with their corresponding buffers, and scattering was measured every 15 min for 6 h.

pH Jump Experiments. Interconversions between N, I_N , I_U , and U were monitored by diluting 10 μ M SMA at one pH into buffer of another pH, such that the final concentration of protein was 1 μ M. After manually mixing, the intensity of either tryptophan fluorescence (excitation 280 nm and emission 345 nm) or ANS fluorescence (excitation 380 nm and emission 470 nm) was monitored, using a time-based scan on a Spex Fluoromax instrument with 1 s time-averaging.

RESULTS

Formation of Amyloid Fibrils Is Favored by Destabilizing Environmental Conditions. The enhanced fluorescence emission of the dye Thioflavin T on association with amyloid fibrils provides a very convenient method to monitor the kinetics of amyloid fibril formation (17, 20). Fibril formation by SMA was investigated by stirring solutions of SMA at various values of pH at 37 °C. At room temperature, or in the absence of stirring, no enhancement of TFT was observed for several days.

The rate of fibril formation from SMA was found to be very sensitive to a number of extrinsic factors, including pH.

agitation, and temperature. As is typical for other amyloid fibril growth curves, SMA fibrillation kinetics exhibit a quasi-sigmoidal behavior, consisting of a lag phase followed by a logarithmic growth phase that eventually plateaus (Figure 1). The slight drop in TFT fluorescence sometimes observed at long time periods may reflect conversion of the mature SMA amyloid fibrils to an alternative fibrillar form that has a lower affinity toward TFT. For example, we frequently observed lateral aggregation of mature fibrils coinciding with the decrease in TFT fluorescence, suggesting a possible decrease in the availability of TFT binding sites.

We confirmed that the enhanced fluorescence emission of TFT was indeed due to interaction with SMA amyloid fibrils by the characteristic Congo red green-birefringence observed under crossed-polarization (data not shown) and by direct observation with transmission electron microscopy and atomic force microscopy. These techniques demonstrated the presence of fibrils in systems with high TFT fluorescence, and their absence in samples with no increase in TFT fluorescence. The TFT assay was shown to be linear over the concentration range of 0 to $>12 \mu\text{g}$ of amyloidogenic protein in the assay solution.

The initial lag in fibril formation (see Figure 1) is often attributed to the slow assembly of a critical nucleus in a nucleation-polymerization mechanism (27). The length of the lag (measured by extrapolating the exponential growth phase to zero intensity) during SMA fibril formation was a sensitive function of the pH at which the fibrils were generated. Some typical data are shown in Figure 1. The length of the lag decreased from days at pH 7.0, 37 °C, for 40 μM SMA, to a few hours at pH 2 (these values for the lag time are very sensitive to the rate of stirring or agitation). In addition, the maximum signal obtained using the TFT assay increased with decreasing pH, indicating that more fibrils were formed at lower pH values (note that the TFT binding assay is performed at pH 7).

Static light scattering was also used to monitor the kinetics of aggregation (Figure 1). Surprisingly, we noted that amorphous aggregation occurred in the same incubation samples of SMA as fibril formation, but at a faster rate. Substantial amorphous aggregation was observed from pH 7 to 4. The amount of amorphous aggregation was proportional to the protein concentration. The amorphous aggregation was observed immediately after starting the stirring at 37 °C as indicated by increased light scattering (Figure 1B,C), whereas the presence of fibrils, as reflected by the increase in TFT fluorescence, was not observed for several days. Confirmation of the fact that this early aggregation was indeed amorphous comes from TEM and AFM micrographs that showed amorphous material and the absence of fibrils (Figure 2). Under certain conditions, e.g., pH 5.0, 37 °C, 100 mM NaCl, 50–60% of the SMA had precipitated (as amorphous aggregate) after 24 h of incubation (based on the absorbance of the supernatant), and no fibrils were visible by microscopy. In contrast, at pH ≤ 3 essentially all of the precipitate was fibrillar within 24 h. The rate of the increase in the light scattering observed at pH 2 correlated with that of the increase in Thioflavin T fluorescence (Figure 1A), suggesting that the predominant species present in solution was fibrillar. This was confirmed by TEM, which indicated that the aggregates were largely fibrillar, though some amorphous material was present. Interestingly the

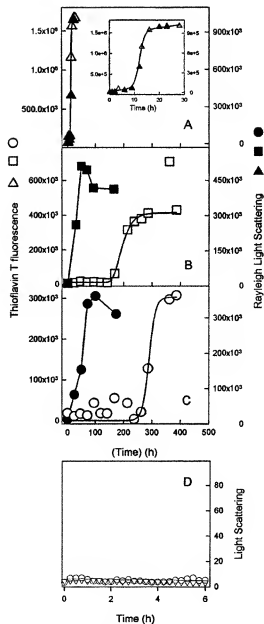


FIGURE 1: pH dependence of amyloid fibril formation by recombinant V_L domain SMA. Fibril formation was monitored using Thioflavin T emission at 482 nm upon excitation at 450 nm at pH 2 (A, Δ), 5 (B, \square), and 7 (C, \circ). Rayleigh light scattering was also monitored at pH 2 (A, Δ), pH 5 (B, \square), and pH 7 (C, \bullet). The formation of amorphous aggregates precedes the formation of amyloid fibrils at pH 7 and 5, conditions that favor the native or the native-like intermediate conformation. The inset to panel A shows an expanded time scale. Light scattering is sensitive to the presence of both amorphous and fibrillar material, whereas TFT fluorescence is selective for fibrillar aggregates alone. Panel D shows that in the absence of agitation, at SMA concentrations as high as 0.5 mM, no aggregation occurs over at least a 6 h period: circles are for pH 7, triangles for pH 2. With agitation, the signal would be >1400 due to the aggregation.

maximum increase in Thioflavin T fluorescence was significantly less at pH 7 compared to pH 5 and pH 2, which also correlated with fewer fibrils observed by microscopy at pH 7 compared to the lower pH conditions.

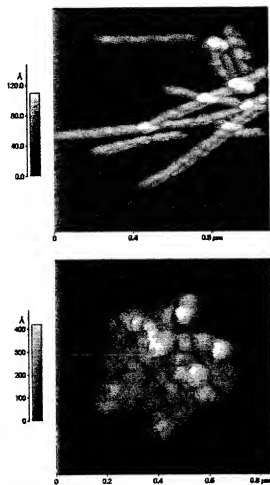


FIGURE 2: Atomic force microscopy images of SMA fibrils observed at pH 2 after 24 h of stirring at 37 °C (A, top). At pH 5 and 7 after incubation for 1 h at 37 °C with constant stirring, a nebulous and loosely packed amorphous deposit is observed (B, bottom). After a few days at pH 5 and 7, both fibrils and amorphous deposits are observed.

Morphology of Fibrillar and Amorphous Deposits. The SMA aggregates were examined using atomic force (Figure 2) and transmission electron microscopy. The images revealed unbranched, rope-like fibrils (Figure 2A), several hundred nanometers in length, most with diameters of ~8 nm, but some, protofibrils, with diameters of ~4 nm (7). Upon incubation at 37 °C at pH 4–6, amorphous aggregates were observed (Figure 2B).

Spectroscopic Characterization of Acidic pH Conformations of SMA. A number of spectroscopic probes were utilized to examine conformational changes in SMA that occur under conditions favoring the formation of amorphous aggregates (pH 4–6) and amyloid fibrils (pH <3). Both amyloid fibrils and amorphous aggregates are only formed in solutions of SMA at 37 °C upon agitation of the solution. Note that all the spectroscopic analyses were done at 37 °C without agitation within 2–3 h of preparation, ensuring that only soluble equilibrium conformations were studied; i.e., neither amorphous nor fibrillar aggregates were present in the spectroscopic analyses. No light scattering was observed for at least 6 h for solutions of SMA at concentrations as high as 7 mg/mL (used in the NMR experiments) at various pH

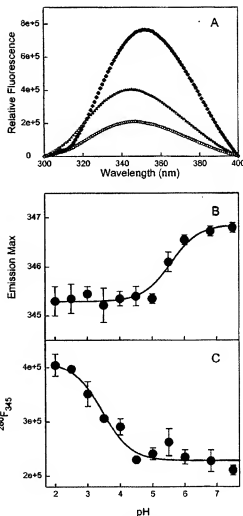


FIGURE 3: Intrinsic tryptophan emission spectra were measured with excitation at 280 nm for 0.5 μ M protein solution in 50 mM buffer and 100 mM NaCl at 37 °C for native SMA at pH 7.5 (\bullet), denatured SMA in 8 M urea at pH 7.5 (\circ), and I_0 at pH 2 (\blacklozenge). The wavelength of maximum emission of tryptophan fluorescence (panel B) and emission intensity (panel C) are plotted against pH. The solid lines are fits to a single ionizable group pH transition using eq 1 (see Materials and Methods). The midpoint of pH transitions for emission maximum and the intensity at 345 nm were 5.6 and 3.5, respectively.

values from 2 to 7 (Figure 1D). Tertiary structure changes were monitored by tryptophan fluorescence emission, near-UV CD, far-UV CD (via the 230 nm peak resulting from aromatic clustering), and by ANS binding studies. Secondary structure changes were monitored by far-UV CD and Fourier transform infrared spectroscopy.

The two tryptophan residues of SMA, W35 and W50, provided convenient spectroscopic means for assessing the protein's conformational state. In particular, W35, which was quenched by the close proximity of the core disulfide formed from C23 and C88 in the native state, was observed to provide a probe of the global conformational state of the V_L domain. Unfolding of the protein resulted in a decrease of the quenching and a consequent increase in the emission intensity of W35 (Figure 3A, and ref 31). The second tryptophan residue, W50, was solvent-exposed in the native

state, based on both the crystal structure of LEN and the observed λ_{max} of 347 nm in native SMA and LEN.

The intrinsic tryptophan fluorescence spectra indicated λ_{max} of 347 and 355 nm for native and denatured SMA (8 M urea), respectively (Figure 3A). In addition, denatured SMA showed a large increase in Trp fluorescence intensity compared to that of its native conformation. When the pH of a solution of SMA was lowered from 7.5, significant changes were observed in the tryptophan fluorescence emission properties. In the pH 4–6 region, there was a decrease in λ_{max} , from 347 to 345 nm (Figure 3B), which was attributed to the build-up of a partially folded intermediate. For reasons to be discussed below, this intermediate was called I_N (for nativelike intermediate). These data were fit to eq 1, and the midpoint of this pH transition was calculated to be 5.6. At pH <4, the emission intensity increases but the λ_{max} does not change further (Figure 3B,C). The increase in emission intensity observed from pH 5 to 2 suggests significant disassembly of the hydrophobic core of the protein. The midpoint of this transition was pH 3.4 and appeared to be cooperative (Figure 3C). The fluorescence spectrum of SMA at pH 2 had a fluorescence intensity between those of the native and denatured states, and a blue shift in the maximum emission to 345 nm. This suggested that substantial residual structure was present at pH 2. This conformation of SMA, populated below pH 3, was called I_U . In addition, the fluorescence spectrum of SMA was measured as a function of protein concentration, over the 0.05–0.5 mg/mL range, to confirm the presence of the two intermediates and to demonstrate the lack of association of the samples under the experimental conditions.

The near-UV CD spectrum of SMA contained significant contributions from the aromatic (tryptophan, tyrosine, and phenylalanine) residues that were sensitive to the tertiary structure of the protein. The near-UV CD spectrum of native SMA (Figure 4A) showed positive peaks at 286 and 296 nm. These likely reflected contributions from two aromatic clusters involving residues Y36, Y86, Y87, F98 and Y27(d), Y32, Y49, Y91, Y92, observed in the crystal structure of LEN which would also be expected to be present in SMA. The peaks at 286 and 296 nm disappeared with transition midpoints of pH 3.2 and 3.4, respectively (Figure 4B). These transitions correspond to the formation of the I_U species, suggesting loss of the nativelike environment of the aromatic groups in this intermediate. The small positive ellipticity for SMA at 268 nm showed transitions with midpoints of 4.9 and 3.7, corresponding to the transitions to I_N and I_U , respectively (Figure 4B). At pH 2, the near-UV CD spectrum of SMA was essentially featureless (Figure 4A), suggesting loss of most of the tertiary structure in I_U , including the aromatic clusters. As a whole, the near-UV CD spectra for SMA in the pH 4–6 region indicated that the underlying tertiary structure is still relatively nativelike in this pH range, consistent with the presence of a nativelike conformation in I_N .

The hydrophobic dye ANS has frequently been used as a probe to reveal the presence of partially folded intermediates due to the presence of increased exposure of contiguous hydrophobic surface area (22–24). ANS did not significantly bind to SMA in its native state, indicating the absence of exposed hydrophobic pockets. However, a pH-titration of SMA in the presence of ANS at 37 °C revealed a marked

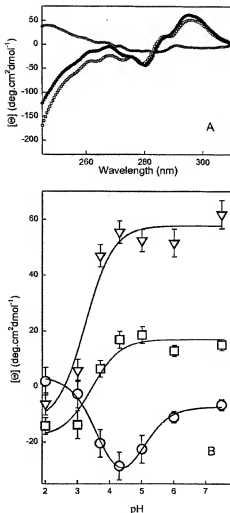


FIGURE 4: Near-UV CD spectra for SMA. Panel A shows plots of molar ellipticity for pH 7.5 (●), pH 5 (□), and pH 2 (Δ). Panel B shows the pH dependence of the molar ellipticity at 296 nm (●), 287 nm (○), and 268 nm (□). The lines through the data are fits to eq 1 for peaks at 296 and 287 nm, yielding apparent pK_a s of 3.2 and 3.4, respectively, which correspond to formation of I_U . The data at 268 nm are fitted to eq 2, resulting in two apparent pK_a s values of pH 4.9 and 3.7. Protein concentration was 0.7 mg/mL.

increase in the fluorescence emission and a blue shift in the ANS emission maximum from 510 to 480 nm. Both the emission intensity increase (data not shown) and the blue shift of the emission were indicative of exposed hydrophobic regions in the vicinity of pH 4–6, with a maximum at pH 4.5 (Figure 5). Such an observation was taken to indicate the build-up of a partially folded intermediate, I_N , which appears to have less contiguous exposed hydrophobic regions. The data suggest that the I_N intermediate is maximally populated between pH 4 and 5 for SMA. In contrast, no ANS binding was observed in the pH 2–10 range for the nonamyloidogenic homologue LEN, indicating the absence of both intermediates with LEN (Figure 5). In addition, the structure of LEN at pH 2 is nativelike, based on small-angle X-ray

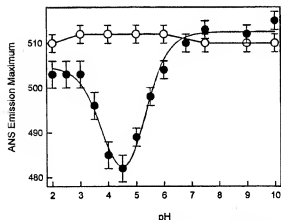


FIGURE 5: pH dependence of binding of ANS to SMA (●) and LEN (○). The reaction was monitored with 0.5 μ M protein solution and 10 μ M ANS, with excitation at 380 nm. Fluorescence emission spectra were collected between 400 and 600 nm at different pHs. The solid line is a fit to two transitions using eq 2 (see Materials and Methods). The midpoints of the pH transitions were 5.2 and 3.8, respectively.

scattering, circular dichroism, FTIR, and Trp fluorescence (unpublished observations).

Circular dichroism spectra were collected for SMA from pH 8 to 2 to probe global secondary structural changes in the different conformers. The far-UV circular dichroism spectrum of native SMA at pH 7.5 was rather unusual, in that it had distinct minima at 230 nm, as well as at 216 nm (Figure 6A). The former was attributed to contributions from aromatic interactions and possibly the disulfide, the latter to β -structure. When the CD spectrum of SMA was examined as a function of pH at 37 $^{\circ}$ C, there were relatively small changes between pH 7.5 and pH 4, with more significant changes occurring at lower pH (Figure 6). The former are consistent with loss of some tertiary structure in I_N . The spectrum at pH 2 was significantly different from the spectrum of SMA denatured in 7 M urea or 5 M Gdn-HCl at pH 7.4, indicating significant structure at pH 2. Plots of the ellipticity at 230 nm against pH reveal the population of a distinct conformational species in the pH 4–6 region (Figure 6B). The ellipticity of SMA monitored at 216 nm (corresponding to β -sheet structure) indicated no change between pH 7.5 and 4.0, suggesting that I_N is relatively nativelylike. However, below pH 4.0, the negative ellipticity at 216 nm shifted toward lower wavelengths, consistent with the loss of β -sheet structure (Figure 6C). The spectrum at pH 2.0 was a mixture of contributions from β -sheet and random coil conformations, indicating some loss of nativelylike β -structure at pH 2. The data were consistent with the presence of a relatively unfolded-like intermediate, I_U , at pH below 3.

Fourier transform infrared spectroscopy (FTIR) has been used to probe protein structure, and the amide I band (1600–1700 cm^{-1}) has been used to estimate protein secondary structure content (25). The ATR-FTIR spectra of hydrated thin films of SMA at pH 7.5 and 5 revealed that significant secondary structural changes occur for the I_N intermediate compared to the native state (Figure 7). The major differences are an increase in low-frequency β -sheet (1625 cm^{-1}), an increase in disordered structure (1648 cm^{-1}), increased turn

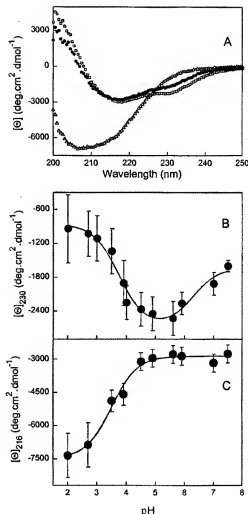


FIGURE 6: Far-UV CD spectra of SMA as a function of pH. Panel A shows the spectra at pH 7.5 (●), pH 5 (□), and pH 2 (Δ). The changes in molar ellipticity at 230 nm and at 216 nm are plotted against pH in panels B and C, respectively. The solid line in panel B is a fit to eq 2 and gives apparent pK_a s of 6.3 and 3.7. The solid line in panel C is a fit to eq 1 and gives an apparent pK_a of 3.5.

(1672 cm^{-1}), and a small decrease in the 1695 cm^{-1} β -component.

At pH 2.0, the amide I spectrum of I_U is different from that at pH 7.5 or 5.0, indicating that I_U has a different secondary structure from native and I_N . The major changes are a large increase in the looplike structure at 1660 cm^{-1} and loss of the major β -peak at 1638 cm^{-1} in the native spectrum, which is replaced by a new, lower intensity β -component at 1631 cm^{-1} .

^1H NMR spectra were collected to further assess the conformational changes that took place in SMA at intermediate and low pH. As shown in Figure 8A, the NMR spectrum of SMA at pH 7.0 is characteristic of a tightly folded protein, having well-dispersed amide, aromatic, and aliphatic proton resonances. As the pH was reduced to below pH 5 (Figure 8B), only minor changes in the spectra were observed. These changes included both sharpening of many resonance lines as well as changes in some amide proton chemical shifts. However, the spectrum was still characteristic of a well-

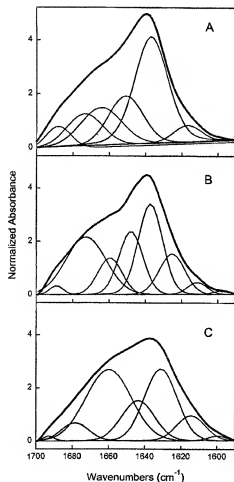


FIGURE 7: Amide I region of the FTIR spectrum of SMA. Panel A shows the spectrum for native SMA at pH 7.5. Panel B shows the spectrum of the nativelylike intermediate (I_N) at pH 5.0, and panel C shows the spectrum of the unfolded-like intermediate (I_U) at pH 2.0. The raw ATR-FTIR spectra after water vapor subtraction are shown as thick lines. The thin lines in each panel are the component spectra obtained after curve-fitting the raw spectra.

folded protein. When the pH was adjusted to below 3, however, the chemical shift dispersion was lost, with the amide proton resonances collapsing to an envelope less than 1.5 ppm wide (Figure 8C). The upfield methyl resonances were also lost below pH 3, with both the aromatic and aliphatic proton regions of the spectrum showing considerable loss of dispersion. Notably, no resonances corresponding to either of the tryptophan indole moieties were apparent in the low-pH spectra. Additionally, the spectrum at pH 2.0 was considerably different from that recorded for SMA under strongly denaturing conditions (pH 2, 8 M urea) as shown in Figure 8D. Under these strongly denaturing conditions, a further loss of dispersion occurs throughout the spectrum, and significant changes in the chemical shifts of nearly all of the amide protons occur. As the solution pH is lowered, the NMR spectra show an increase in signal-to-noise for the same concentration of protein. This is likely due to dissociation of the V_1 dimer ($K_d = 40 \mu\text{M}$ at pH 7) at lower pH.

Small-Angle X-ray Scattering Characterization of SMA Conformations. Small-angle X-ray scattering measurements indicated that SMA became less compact as the pH was

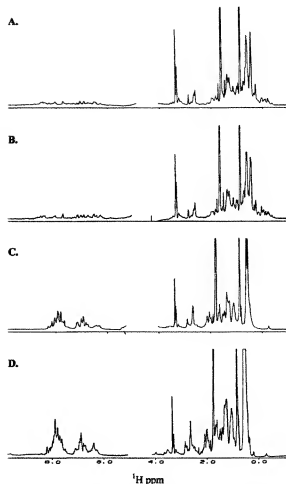


FIGURE 8: ^1H NMR spectra of SMA. Panel A shows the spectrum of the native protein at pH 7. Panel B shows the spectrum of the nativelylike intermediate, I_N , at pH 5, and panel C shows the spectrum of the relatively unfolded intermediate, I_U , at pH 2. For comparison, the spectrum of the unfolded protein in 8 M urea, pH 2, is shown in panel D.

reduced from 7 to 2 (at a protein concentration of $80 \mu\text{M}$), with an increase in R_g from $19.6 \pm 0.4 \text{ \AA}$ at pH 7 to $26.8 \pm 0.6 \text{ \AA}$ at pH 2. At pH 5, the protein was only slightly expanded, having a R_g of $20.5 \pm 0.6 \text{ \AA}$. Kratky plots of the scattering data indicated that extensive globularity was maintained even at low pH, although some denaturation was apparent at pH 5 and more so at pH 2. The significant compactness of I_U ($R_g = 26.8 \text{ \AA}$) is apparent by comparison of its R_g to that of the fully unfolded protein ($R_g > 30 \text{ \AA}$).

Equilibrium Thermodynamic Stability of SMA. The stability of SMA was measured at different pHs using urea denaturation monitored by intrinsic tryptophan fluorescence and far-UV CD. Tryptophan fluorescence had the advantage that it permitted the use of low protein concentrations, which eliminated potential aggregation problems during unfolding. At pH 7.5, SMA is only marginally stable, with a free energy of unfolding $\Delta G^\circ = 4.8 \text{ kcal/mol}$ and $m = 1.05 \text{ kcal/mol}$. Similar equilibrium plots were obtained when starting with either native or unfolded protein, indicating that there was no hysteresis in the urea unfolding transitions. As the pH was decreased, the stability of SMA decreased significantly (Table 1) as indicated by the decrease in the midpoint of

Table 1: Effect of pH on the Stability of SMA*

pH	C_{50} (M)
2	~0
4	1.5 \pm 1.5
6	3.2 \pm 0.1
8	4.0 \pm 0.1
10	3.5 \pm 0.1

* Midpoints of urea unfolding transitions were monitored by Trp fluorescence.

the urea unfolding transition with decrease in pH. In light of the evidence for formation of a non-native species from SMA in the vicinity of pH 4–6 (I_N), the urea unfolding data were not converted into free energy data except for neutral pH.

Kinetics of Interconversion of Intermediates. The rates of interconversion between the native conformation (N) and the partially folded intermediates (I_N and I_U) were monitored by pH jumps. Interconversions between N and I_N were monitored using changes in the ANS fluorescence and jumping the pH of solutions of SMA from 7 to 5 (N to I_N) and from 5 to 7 (I_N to N). Interconversions involving I_U were followed by observing changes in the tryptophan fluorescence, with pH jumps of 7 to 2 (N to I_U), 5 to 2 (I_N to I_U), 2 to 5 (I_U to I_N), and 2 to 7 (I_U to N). The results from these experiments show that conversion of N to I_N and I_N to N are fast processes, complete within the dead time of manual mixing, indicating that the rate constant is >0.35 s⁻¹. On the other hand, conversion of either N or I_N to I_U and the reverse are slower processes. The rates for both N to I_U and I_N to I_U were the same within experimental error, namely, 0.01 ± 0.002 s⁻¹, consistent with I_N lying on the pathway between N and I_U . The rates of the transformation of I_U to either I_N or N were the same, with a rate constant of 0.002 ± 0.0008 s⁻¹, again consistent with I_N lying on the pathway between N and I_U .

DISCUSSION

There is increasing evidence to suggest that protein aggregation, including amyloid fibril formation, arises from a partially folded conformation of the aggregating protein (5, 6, 26–30). The present data strongly support such a hypothesis for the aggregation of immunoglobulin light chain variable domains. Protein aggregation has generally been regarded as being driven by nonspecific hydrophobic interactions operating on unfolded or collapsed molten-globule states. On the other hand, extracellular aggregation of some proteins to form amyloid fibrils has been conventionally attributed to mutations altering the local surface properties of the native state, thereby introducing new packing interactions for oligomerization of the native state (26, 31). In the case of SMA, our results clearly indicate that a nativelike conformation in the fibrils is highly unlikely. Thus, the model for light chain amyloid fibrils proposed by Stevens and co-workers (31), and based on the native structure, is inconsistent with the experimental observations.

The partially folded conformations that are the critical precursors to protein aggregation may arise either during the folding of newly synthesized proteins, as with inclusion bodies, or from the native state, as appears plausible for the extracellular amyloid deposits. The build-up of the soluble

precursor that triggers aggregation could involve a combination of factors including an amino acid sequence leading to a relatively less stable native state as compared to that of a nonaggregating variant and/or the presence of mildly destabilizing extrinsic conditions.

Detailed analysis of the properties of the amyloidogenic SMA, including its stability, conditions necessary to populate partially folded intermediate conformations, propensity to aggregate, and kinetics of aggregation and fibril formation, provides insight into the molecular basis for aggregation. The present results raise a number of interesting questions, such as: Which features are responsible for the propensity to form fibrils? Why are both amorphous and fibrillar aggregates formed, and what is the relationship, if any, between them? How do the two partially folded intermediate conformations, I_N and I_U , fit into the picture? We will begin with this last question.

Destabilizing Conditions Lead to Partially Folded Intermediate Conformations. Mildly destabilizing conditions, such as low pH or low urea concentrations (data not shown), lead to enhanced aggregation and fibrillation of SMA. The results of the spectroscopic investigations of SMA as a function of pH reveal that SMA forms two partially folded conformations, I_N and I_U , the former being relatively nativelike in its structural properties, whereas the latter is considerably more unfolded. The major significance of the observation of these species is the correlation between formation of amorphous aggregates and I_N , and formation of fibrils and I_U . Both I_N and I_U are envisaged as ensembles of conformations that retain some nativelike structure (more in I_N and less in I_U) with the remainder of the chain, especially for I_U , being highly mobile, and disordered but biased toward its native conformation.

The near- and far-UV circular dichroism, Trp fluorescence, NMR, and SAXS data all point to I_N as being a relatively nativelike species with most structural properties similar to those of the native state. The significant increase in ANS binding, however, points to the critical feature of this intermediate, namely, increased exposure of hydrophobic surfaces compared to the native state. The increased negative ellipticity in the far-UV CD at 230 nm is related to this as it probably represents minor structural rearrangements in side-chain packing manifested as changes in the CD contribution of aromatic residues, rather than secondary structure changes. The enhanced ANS binding in the pH 4–6 range is very consistent with the population of a partially folded intermediate (22, 23). Likewise, the FTIR spectrum for I_N reveals that although the overall secondary structure is quite similar to that of the native state, nevertheless there are significant structural differences. These include increased turn structure and a shift in some of the β -structure to components with a lower frequency band, perhaps signifying changes in the β -strand interactions. From examination of the spectral probes as a function of pH, it is apparent that there is a structural transition with a midpoint around pH 5.5. This transition, which corresponds to the interconversion of the native state to I_N , could reflect the titration of histidine or carboxylate residues in SMA. The pH-dependent transition from N to I_N was not observed for the nonamyloidogenic LEN (Figure 5 and unpublished observations). The only differences in ionizable side chains between LEN and SMA are two histidines present in SMA (8); this suggests that one

or both of these His residues, either directly or indirectly, may be responsible for the transitions between N and I_N.

The data indicate that SMA forms a second, more unfolded, intermediate, I_U, at pH <3. From comparison of the spectral probe signals as a function of pH, a common transition, attributed to that of I_N to I_U, with a midpoint in the vicinity of pH 3.3 is observed with all the probes. This intermediate retains substantial compactness, and secondary structure, consistent with the presence of a partially folded intermediate conformation. Based on the apparent pK of the transition between I_N and I_U, this conformational change is apparently governed by the titration of carboxylate groups. The fact that at pH 2 the fluorescence signal reflects substantial residual structure (significantly decreased intensity and blue-shifted λ_{max} relative to the unfolded and native states) suggests that there is also tertiary structure present in I_U. However, the near-UV CD spectrum indicates the loss of most of the native aromatic side-chain interactions, suggesting that the aromatic clusters present in the native conformation may no longer be present. The FTIR spectra indicate additional loop/disordered structure in I_U compared to the native and I_N conformations and also loss of nativelike β -structure. The NMR spectrum of this intermediate also clearly indicates that it retains considerable secondary and tertiary structure.

Among the factors that stimulate fibril formation from SMA are increased protein concentration, agitation, and increased temperature. All these are likely to result in increased concentration of non-native conformations (through equilibrium between native and non-native states, denaturation at air-water interfaces, and shifting of the equilibrium from native to non-native conformations, respectively), with their known propensity to aggregate. These observations strengthen the correlation between aggregation and precursor partially folded intermediates.

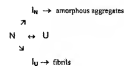
A minor complication in some of these experiments arises from the fact that V₁ domains are known to dimerize in a fashion similar to the association of the C₁/V₁ domain interaction in intact immunoglobulins. Stevens and Schiffer (32) demonstrated that native SMA exists in a monomer-dimer equilibrium with $K_d = 40 \mu\text{M}$ under physiological conditions. Thus, data collected with high concentrations of SMA, such as SAXS and NMR, will be potentially complicated by the presence of these native dimers, at least at neutral pH. The main anticipated effect would be that under such conditions the equilibrium between the native conformation and the intermediates would be shifted in favor of the native conformation, rather than a non-native one. This leads to a potential decrease in the rate of fibril formation under conditions where the protein is present mostly as the native dimer (unpublished observations).

Relation between Stability and Aggregation. The stability of SMA is greatly decreased at acidic pH, correlating with both increased amorphous and fibrillar aggregation. The fact that at lower pH SMA readily forms fibrils suggests that removal of nativelike interactions is important prior to fibrillation. Our data show that the aggregation of the amyloidogenic SMA correlates well with the decreased stability of the native state and the population of non-native conformations. We believe that it is the *differential destabilization* of the native state of SMA relative to the partially folded intermediate conformations which is the key feature

Scheme 1



Scheme 2



of the amyloidogenesis, and which will be attenuated in nonamyloidogenic light chains. These observations are in accord with previous investigations of the correlation between amyloidogenesis and stability in light chains, which has indicated that either destabilizing mutations or sequences (12, 33, 34), or destabilizing conditions (35) correlate with increased fibrillogenicity. In fact, SMA has been shown to be about 2.5 kcal/molecule less stable than its benign LEN homologue (8). Further, it has been shown that destabilizing LEN with Gdn-HCl leads to fibrillation (12).

Amorphous and Fibrillar Aggregates. The observation that both fibril and amorphous aggregate formation occur simultaneously under many conditions raises a number of questions, for example: Do they arise from similar or different partially folded conformations, and do amorphous aggregates convert directly or indirectly to fibrils?

The coincident kinetics for light scattering and TFT at pH ≤ 3 , in conjunction with the limited amount of amorphous aggregates observed by EM, indicate that at these low pHs fibrils are preferentially formed, and that the limited amount of amorphous aggregates formed under these conditions rapidly converts to fibrillar species. Since the spectroscopic results suggest that at pH ≤ 3 the only significant species present is I_U, the limited amount of amorphous aggregates indicates that it is I_U that is responsible for fibril formation. Similarly, the fact that the pH at which maximal amounts of amorphous material is found is in the vicinity of 5.5 indicates that I_N is primarily responsible for the amorphous aggregates. The decreased amplitude of the final TFT signal at higher values of pH is attributed to the increased amount of amorphous aggregate at higher pHs.

The kinetics of interconversion between N, I_N, and I_U are consistent with I_N being on the pathway between N and I_U (Scheme 1). However, the possibility that there are separate pathways to I_N and I_U (Scheme 2) cannot be eliminated at this time. The fast interchange between N and I_N is not surprising, given the fact that I_N is relatively nativelike.

The decreased amounts of fibrils observed at higher pH values presumably result from the smaller amounts of I_U in equilibrium with N and I_N (even though the equilibrium levels of I_U may be quite low at higher pHs, any I_U lost in fibril formation will be replaced by mass action with more soluble I_U). The strong correlation between the pH dependence of aggregation (and fibril formation) and the population of the partially folded intermediates supports the hypothesis that the observed intermediates are key players in the aggregation process. Fibril generation, even under nativelike conditions, can be attributed to the Boltzmann distribution of ensembles of various states under nativelike conditions. Hence, it is possible that the key intermediate, highly populated

at pH ≤ 3 , is also present under nativelike conditions, but at substantially lower concentration, and is responsible for amyloid formation after an extended lag period.

Aggregation results from the strong self-association tendency of the partially folded intermediates, probably due to the presence of large solvent-exposed hydrophobic patches, which are absent in the native and fully unfolded states. The increased β -structure observed in the aggregated states reflects β -strand- β -strand interactions involved in the intermolecular association. Fibril formation is expected to involve a number of intermediate states of soluble oligomers of partially folded intermediates, potentially populated at very low levels. Aggregation occurs under conditions in which a suitably high concentration of the key partially folded intermediate is present, due to a combination of destabilizing factors, such as pH and temperature, or urea, and amino acid sequence, as well as the concentration of the intermediate. Thus, it is mostly the intrinsically low stability of SMA that leads to the build-up of the intermediate leading to aggregation, under conditions where more stable light chains form negligible intermediate and remain in the native conformation.

The much more rapid formation of amorphous aggregates of SMA, compared to fibrils, is very interesting, and raises a number of questions regarding the nature of the relationship between the initially formed amorphous aggregates and the more slowly formed fibrils. Based on the data reported here, it is apparent that the partially folded intermediate populated in the pH 4–6 region is the direct precursor of the amorphous aggregates. The correlation between I_A and amorphous aggregates, and I_H and fibrillar aggregates, suggests that the ratio of the two types of deposits is determined, at least in part, by kinetic competition between the pathways leading to the two different intermediates. A more detailed investigation of the relationship between amorphous and fibrillar deposits will be given elsewhere.

The results of the present investigation firmly establish the existence of partially folded intermediates as key precursors on the aggregation pathway of the amyloidogenic light chain variable domain SMA. In addition, the observation of two such intermediates is the first report that a given protein might have more than one critical intermediate conformation on the aggregation pathway, and that such different conformations may lead to different types of deposits. One implication of this is that factors, such as chaperones, which may change the effective concentration of one of the intermediates may change the nature of the deposits.

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parts of a cell. In addition to compartmentation, cellular processes are regulated by enzyme synthesis and destruction. Often, enzymes are synthesized at low rates until increased demands by the cell (for instance, appearance of substrate), new enzymes are synthesized. Later, the pool of enzyme is lowered when levels of substrate decrease or the cell becomes inactive.

SUMMARY Functional Design of Proteins

The function of nearly all proteins depends on their ability to bind other molecules (ligands). Ligand-binding sites on proteins and the corresponding ligands are chemically and topologically complementary. The strength of binding is specifically, to the restriction of binding to one or a few preferred ligands.

Enzymes are catalytic proteins that accelerate the rate of cellular reactions by lowering the activation energy and stabilizing transition-state intermediates.

Enzyme active site consists of a catalytic region, a binding site, and an allosteric region. The amino acids composing the active site are not necessarily adjacent in the amino acid sequence, but are brought into proximity in the native conformation.

- The kinetics of many enzymes are described by the Michaelis-Menten equation. The kinetic parameters of an enzyme can be determined: the Michaelis constant K_m is a measure of the enzyme's affinity for substrate, and the maximal velocity V_{max} (see Figure 3.2b).
- Many allosteric enzymes are regulated by a cofactor, a molecule in substrate, activator, or inhibitor induces a conformational change, or allosteric transition, that alters the protein's activity or affinity for other ligands.
- In multimeric proteins that bind multiple ligands, the binding of one ligand to one subunit influences the binding of subsequent ligand molecules. Enzymes that cooperatively bind substrates exhibit sigmoidal kinetics (see Figure 3.2b).

Allosteric mechanisms can act like switches, turning protein activity on or off. Covalent modification is another means of regulation. Phosphorylation can have the same regulatory effect. Phosphorylation is reversible; converts inactive zymogens into active enzymes.

3.4 Membrane Proteins

As we've seen, all antibodies have a similar structure and function; enzymes are structurally varied, but all have a catalytic function. In contrast, although all membrane proteins

are located at the membrane, they otherwise are both structurally and functionally diverse. In this section, we describe the membrane as a fluid mosaic. The fluid mosaic model of the membrane has the same basic phospholipid bilayer structure. Associated with each membrane is a set of membrane proteins that enable the membrane to carry out its functions (see Figure 3.3). The composition of the proteins attached to a membrane varies depending on cell type and subcellular location.

Some proteins are bound only to the membrane surface, and others are embedded in the membrane. Proteins located on one or both sides of it. Protein domains on the extracellular membrane surface are generally involved in cell-cell signaling or interactions. Domains within the membrane, particularly those that form transmembrane α helices, are involved in signal transduction. Domains lying along the cytosolic face of the membrane have a wide range of functions, from anchoring cytoskeletal proteins to the membrane to regulating intracellular signaling pathways. In many cases, proteins are associated with the membrane by covalently attached polypeptide chains in the membrane can be predicted based on its homology with another, well-characterized protein. In this section, we examine the characteristic structural features of membrane proteins and then describe the structure and function of various types of membrane proteins as presented in several later chapters. The synthesis and processing of membrane proteins are discussed in Chapter 17.

Proteins Interact with Membranes

In different ways, proteins can be classified into two broad categories: integral and peripheral. Integral proteins are embedded in the membrane, interacting with the lipid bilayer on the nature of the membrane-protein interactions (see Figure 3.3). Most biomembranes contain both types of membrane proteins.

Some membrane proteins, also called *intrinsic* proteins, have one or more segments that are embedded in the phospholipid bilayer. Most integral proteins contain residues with hydrophobic side chains that interact with the hydrophobic core of the membrane, anchoring the protein to the membrane. Most integral proteins span the entire phospholipid bilayer. These *transmembrane* proteins contain one or more membrane-spanning domains as well as domains in the aqueous medium on each side of the bilayer. In the transmembrane proteins examined to date, β strands in contrast, soluble proteins are predominantly bound to the membrane by noncovalent interactions. The polypeptide acid, as discussed later, in these proteins, the bound fatty chain does not enter the phospholipid bilayer. Proteins, do not interact with the hydrophobic core of the phospholipid

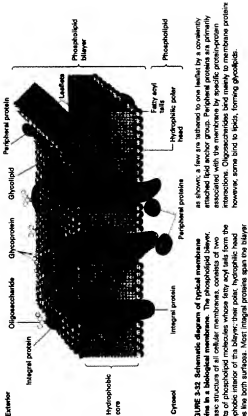


FIGURE 3.3 Schematic diagram of typical membrane structure in a biological membrane. The composition of the basic structure of all cellular membranes consists of two layers of phospholipid molecules whose fatty acid tails form the hydrophobic core of the membrane. The hydrophilic heads of the phospholipids are exposed to the aqueous environment on both sides of the bilayer.

bilayer. Instead they are usually bound to the membrane indirectly by interactions with integral membrane proteins or directly by interactions with lipid polar head groups. Membrane proteins that are not embedded in the lipid bilayer include the cytoskeletal proteins spectrin and ankyrin in erythrocytes (Chapter 18) and the enzyme protein kinase C. The enzyme ankyrin between the cytosol and the cytosolic face of the membrane (see Figure 3.3). Ankyrin plays a role in signal transduction (see Chapter 20). Other non-covalently bound proteins (see Figure 3.3) are associated with the membrane by certain proteins of the extracellular matrix, are localized to the outer (epithelial) surface of the plasma membrane.

Hydrophobic α Helices in Transmembrane Proteins Are Embedded in the Bilayer

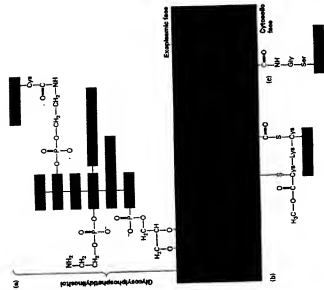
Integral proteins containing membrane-spanning α helices are embedded in the membrane. The hydrophobic α helices interact with the lipid interior of the bilayer and probably also by ionic interactions with the polar head groups of the phospholipids. Glycophorin, a major erythrocyte membrane protein, is a single-pass transmembrane protein (see Figure 3.3). Glycophorin contains a membrane-spanning α helix composed entirely of hydrophobic (or uncharged) amino acids. The predicted length of this α helix (27.5 nm) is just sufficient to span the hydrocarbon core of the bilayer (see Figure 3.3). The hydrophobic α helices shield the polar carbonyl ($C=O$) and the amino (NH) groups of

the peptide bond, which are all hydrogen-bonded to one another. This hydrophobic helix is prevented from slipping across the membrane by a flanking set of positively charged residues. The hydrophobic α helices are associated with negatively charged phospholipid head groups. In glycoproteins, most of these charged residues lie adjacent to the cytosolic helix.

Many Integral Proteins Contain Multiple Transmembrane α Helices

Although Figure 3.3 depicts glycophorin as a monomeric integral protein, many integral proteins are dimers. This is present in erythrocyte membranes as a dimer of two identical polypeptide chains. The two membrane-spanning α helices of glycophorin are thought to form a coiled-coil structure between the amino acid side chains at the interface of the two helices. It is now known that many other transmembrane proteins contain two or more membrane-spanning α helices. For example, the β -subunit of the Na^+ /K⁺ ATPase (see Figure 3.3) contains four α helices and several protein groups, including four chlorophyll molecules. In the complex protein, three of the four subunits span the membrane, two of these subunits (L and M) contain a transmembrane α helix. A large and important family of integral proteins is defined by the presence of seven membrane-spanning α helices.

4 FIGURE 3-38 Anchoring of integral proteins to the plasma membrane by hydrophobic and covalent bonds. (a) The protein and several hydrophobic enzymes are anchored to the phospholipid bilayer by hydrophobic interactions. This is the case for the cytochrome *b₅* complex. (b) Cytochrome *c* is anchored to the cytosolic leaf of the membrane through transfer and covalent bonds. (c) The cytochrome *b₅* complex is anchored to the membrane through myristate and another fatty acid attached to an N-terminal glycine residue.



including alkaline phosphatase, fall into this class. Various experiments have shown that the phospholipid anchor is both necessary and sufficient for limiting the mobility of these enzymes. For instance, the enzyme phospholipase C cleaves the phosphate-glycerol bond in phospholipids as well as in glycerophosphatidylcholine (GPC) anchors, and treatment of cells with phospholipase C causes the release of enzymes such as alkaline phosphatase and Thy-1 protein and alkaline phosphatase from the cell surface.

Some cytosolic proteins are anchored to the cytoplasmic leaf of the membrane by covalent bonds. The most common anchors are prenyl, farnesyl, and geranylgeranyl groups. These proteins undergo a chemical modification involving the attachment of a prenyl group to the C-terminus of the protein. The modified protein then undergoes proteolysis and methylation; these reactions remove the three terminal amino acids and the prenyl group, leaving a C-terminus. In these cases, fatty acyl palmitate groups

from thioester bonds to nearby cysteine residues, providing additional anchors and limiting mobility. The myristate anchor is common to the membrane (see Figure 3-36b). In another group of lipid-anchored cytosolic proteins, a fatty acid group (e.g., myristate or palmitate) is linked by an amide bond to the N-terminus of the protein. This is necessary for retention at the membrane and may play an important role in a membrane-associated function. For example, $\gamma\text{-Sec}$, a mutant form of $\gamma\text{-glutamyl transcarboxylase}$ in which the N-terminus of the enzyme is myristoylated (N-termallyl) Neomycin.

Some Peripheral Proteins Are Soluble Enzymes That Act on Membrane Components

An important group of peripheral membrane proteins are water-soluble enzymes that associate with the membrane by a hydrophobic interaction with the hydrophobic group of such enzymes and the phospholipids, which hydrolyze various bonds in the head groups of phospholipids



A FIGURE 3-37 Schematic diagram of a phospholipid bilayer. The head group of a phospholipid is attached to the protein in the outer leaflet. The head group of a phospholipid is attached to the protein in the inner leaflet. The head group of a phospholipid is attached to the protein in the inner leaflet.

(Figure 3-37). These enzymes have an important role in the degradation of damaged or aged cell membranes. For example, phospholipase A₂ and B₂ are enzymes that hydrolyze the ester bonds in phospholipids, releasing fatty acids and glycerol. These enzymes are found in many tissues and are involved in a variety of biological processes. For example, phospholipase A₂ is involved in the regulation of blood clotting and in the release of arachidonic acid from membrane phospholipids. Phospholipase B₂ is involved in the regulation of cell growth and differentiation.

SUMMARY Membrane Proteins

1. Biological membranes usually consist of both integral and peripheral membrane proteins (see Figure 3-32).
2. Integral membrane proteins include transmembrane proteins and lipid-anchored proteins.
3. Two types of membrane-spanning domains are found in transmembrane proteins: one or more α helices or β barrels.
4. Some integral membrane proteins are associated with the membrane by covalent bonds to the lipid head groups, while others are associated by noncovalent interactions with the lipid head groups.
5. When the polypeptide chain of a transmembrane protein spans the lipid bilayer, the core of the protein is hydrophobic and is surrounded by the hydrophobic tails of the phospholipids.
6. Some integral membrane proteins are associated with the membrane by covalent bonds to the lipid head groups, while others are associated by noncovalent interactions with the lipid head groups.

- Amino acid residues modified with long-chain hydrocarbon and/or other nonpolar groups are common to many integral proteins to form a membrane phospholipid.
- Peripheral membrane proteins interact with integral membrane phospholipids or with the polar head groups of membrane phospholipids. They do not enter the hydrophobic core of the membrane.

3.5 Purifying, Detecting, and Characterizing Proteins

A protein must be purified before its structure and the mechanisms of its action can be studied. However, because proteins are complex molecules, their purification is a difficult task. In this section, we describe some of the methods that can be used to isolate a protein. To isolate one particular protein from the estimated 10,000 different proteins in a cell is a daunting task that requires methods both for separating proteins and for detecting the presence of specific proteins.

Any molecule, whether it is a protein or a small molecule, can be separated from other molecules based on large differences in some physical characteristic. Although the separation of amino acids is a process that is usually performed in the laboratory, the same principles can be applied to the separation of proteins. In this section, we briefly outline different techniques for separating proteins based on their size and other properties. Transferring proteins based on their size and other properties is a common technique for separating proteins. We then consider general methods for detecting or assaying specific proteins, including the use of radioactive compounds for tracking biological activity. Finally, we describe some of the methods for determining a protein's mass, sequence, and three-dimensional structure.

Proteins Can Be Removed from Membranes by Detergents or High-Salt Solutions

Because water-soluble globular proteins have many exposed hydrophilic groups, they maintain their native conformation in aqueous solution. However, when they are removed from membranes, they lose their native conformation. To isolate a protein from a membrane, the proteins are separated from the membrane, their exposed hydrophobic regions interact, causing the protein molecules to aggregate. This process is called denaturation. Denaturation can be achieved by detergents, which have affinity both for hydrophilic groups and for water.

Detergents are amphipathic molecules that disrupt membranes by interacting with the hydrophobic tails of the phospholipids and the hydrophilic heads of the phospholipids. When a detergent is added to a membrane, it disrupts the membrane by interacting with the hydrophobic tails of the phospholipids and the hydrophilic heads of the phospholipids. This process is called denaturation. Denaturation can be achieved by detergents, which have affinity both for hydrophilic groups and for water.